

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 16:01:23 ; Search time 735.656 Seconds
(without alignments)
1383.200 Million cell updates/sec

Title: US-10-729-421-40

Perfect score: 21

Sequence: 1 cagtgcacatgcagggttagct 21

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 100 summaries

Database :

GenEmbl:*
1: gb_ba:*
2: gb_hgt:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vi:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	19.4	92.4	232064	2	AC109544
C 2	19.4	92.4	258823	2	AC133226
C 3	18.4	87.6	140394	2	CR352267
C 4	18.4	87.6	155858	2	AP000772
C 5	18.4	87.6	161516	2	CR391906
C 6	17.8	84.8	40392	3	U21308
C 7	17.8	84.8	72655	2	AC100344
C 8	17.8	84.8	116076	10	AL831718
C 9	17.8	84.8	125354	2	AC148761
C 10	17.8	84.8	139894	10	AC110379
C 11	17.8	84.8	144142	10	AC102916
C 12	17.8	84.8	162289	4	AC097230
C 13	17.8	84.8	162560	2	AC069005
C 14	17.8	84.8	167878	9	AC103719
C 15	17.8	84.8	189662	9	AC015468
C 16	17.8	84.8	197796	2	AC129792
C 17	17.8	84.8	197796	2	AC129792
C 18	17.8	84.8	203690	2	AC087221
C 19	17.8	84.8	214765	10	AC115746

C 20	17.8	84.8	222540	2	AC120123	AC120123 Mus muscu
C 21	17.8	84.8	260600	2	AC115307	AC115307 Rattus no
C 22	17.8	84.8	281447	2	AC129380	AC129380 Rattus no
C 23	17.4	82.9	5446	6	BD185177	BD185177 Novel gen
C 24	17.4	82.9	46070	2	AC121561	AC121561 Homo sapi
C 25	17.4	82.9	85566	9	ALA33227	ALA33227 Human DNA
C 26	17.4	82.9	89171	9	AC008404	AC008404 Homo sapi
C 27	17.4	82.9	92946	2	AC140020_3	Continuation (4 of
C 28	17.4	82.9	93714	2	ALI161661	ALI161661 Homo sapi
C 29	17.4	82.9	110000	2	AC139803_1	Continuation (2 of
C 30	17.4	82.9	118499	9	AC121562	AC121562 Homo sapi
C 31	17.4	82.9	134878	9	AC140847	AC140847 Homo sapi
C 32	17.4	82.9	146174	2	AC138823	AC138823 Homo sapi
C 33	17.4	82.9	146597	2	AC121323	AC121323 Homo sapi
C 34	17.4	82.9	154957	2	AC080126	AC080126 Homo sapi
C 35	17.4	82.9	159347	2	AC138971	AC138971 Homo sapi
C 36	17.4	82.9	170361	2	AC008542	AC008542 Homo sapi
C 37	17.4	82.9	171398	2	AC141597	AC141597 Homo sapi
C 38	17.4	82.9	172276	9	AC139795	AC139795 Homo sapi
C 39	17.4	82.9	174164	9	AC116166	AC116166 Homo sapi
C 40	17.4	82.9	175612	2	AC141254	AC141254 Homo sapi
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C 53	17.4	82.9	197972	9	AC138865	AC138865 Homo sapi
C 54	17.4	82.9	201638	2	AC144987	AC144987 Homo sapi
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C 65	17	81.0	192355	10	AC124126	AC124126 Mus muscu
C 66	17	81.0	196835	2	AC142538	AC142538 Homo sapi
C 67	17	81.0	200000	2	AC004630	AC004630 Homo sapi
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C 69	17	81.0	206833	10	AC124178	AC124178 Mus muscu
C 70	17	81.0	207436	2	AC140823	AC140823 Homo sapi
C 71	16.8	80.0	1265	10	BC016537	BC016537 Mus muscu
C 72	16.8	80.0	1576	5	BX950346	BX950346 Gallus ga
C 73	16.8	80.0	4952	1	MVHUGAUB	X61204 M.voltae va
C 74	16.8	80.0	9431	9	BX571808	BX571808 Human DNA
C 75	16.8	80.0	293364	3	CXC2787	Z54236 Caenorhabdi
C 76	16.8	80.0	39872	9	HSICB2046	Z97183 Human DNA
C 77	16.8	80.0	69184	9	CR759793	CR759793 Human DNA
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C 81	16.8	80.0	79175	2	AC1012516	AC1012516 Homo sapi
C 82	16.8	80.0	83962	2	AP000448	AP000448 Homo sapi
C 83	16.8	80.0	85975	9	AC110011	AC110011 Homo sapi
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C 88	16.8	80.0	110000	2	AC129386_1	Continuation (2 of
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C 90	16.8	80.0	110794	9	AL662827	AL662827 Human DNA
C 91	16.8	80.0	112209	10	AC022298	AC022298 Mus muscu
C 92	16.8	80.0	114575	9	BX248088	BX248088 Human DNA

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93 16.8 80.0 116695 2 CR762434
94 16.8 80.0 117417 9 AL353704
95 16.8 80.0 128293 10 AL928607
96 16.8 80.0 139807 2 AC105325
C 97 16.8 80.0 153700 2 AC118114
C 98 80.0 159703 10 AC122439
99 16.8 80.0 163758 2 AC141345
100 16.8 80.0 166305 2 AC022580

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ALIGNMENTS

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RESULT 1
AC109544
LOCUS
DEFINITION
Rattus norvegicus clone CH230-202010, *** SEQUENCING IN PROGRESS

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ACCESSION
AC109544
VERSION
HTG; HTGS_PHASE2; HTGS_DRAFT; HTGS_ENRICHED.
KEYWORDS
Rattus norvegicus (Norway rat)
SOURCE
Rattus norvegicus
ORGANISM
Rattus norvegicus

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Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

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REFERENCE

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AUTHORS
Muzny,D,Marie, Allen,C, Allen,H, Aylsworth,S, Amin,A, Anguiano,D,
Anyalebechi,V, Aoyagi,A, Ayodeji,M, Baca,E, Baden,H,
Baldwin,D, Bandaranaike,D, Barber,M, Barnstead,M, Benahmed,F,
Biswal,K, Blair,J, Blankenburg,K, Blyth,P, Brown,M,
Bryant,N, Buhay,C, Burch,P, Burrell,K, Calderon,E,
Cardenas,J, Carter,K, Cavazos,I, Ceasar,H, Center,A,
Chacko,J, Chavez,D, Chen,G, Chen,R, Chen,Y, Chen,Z, Chu,J,
Cleveland,C, Cockrell,R, Cox,C, Coyle,M, Cree,A, D'Souza,L,
Davila,M,L, Davis,C, Davy-Carroll,L, De Anda,C, Dederich,D,
Delgado,O, Denison,S, Deramo,C, Ding,Y, Dinh,H, Divya,K,
Draper,H, Dugan-Rocha,S, Dunn,A, Durbin,K, Duval,B, Eaves,K,
Egan,A, Escotto,M, Eugene,C, Evans,C,A, Falls,T, Fan,G,
Fernandez,S, Finley,M, Flagg,N, Forbes,L, Foster,M, Foster,P,
Fraser,C,M, Gabisi,A, Ganta,R, Garcia,A, Garner,T, Garza,M,
Ganeshkumar,L, Gargueta,M, Gargueta,M, Gargueta,M, Gargueta,M,
Gharat,P, Haaland,P, Haines,A, Henderson,N, Hernandez,J,
Harvey,Y, Havlak,P, Hawes,A, Hamilton,C, Hamilton,K,
Hernandez,R, Hines,S, Hladik,S,L, Hodgson,A, Hogues,M,
Hollins,B, Howells,S, Hulyk,S, Hume,J, Idlebird,D, Jackson,A,
Jackson,L, Jacob,L, Jiang,H, Johnson,B, Johnson,R, Jolivet,A,
Karpatis,S, Kelly,S, Kelly,S, Khan,Z, King,L, Kovar,C,
Kowis,C, Kraft,C,L, Lebow,H, Levan,J, Lewis,L, Li,Z, Liu,J,
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Maheshwari,M, Mahindartne,M, Mahmoud,M, Mallory,K, Mangum,A,
Mangum,B, Mapua,P, Martin,K, Martin,R, Martinez,E,
Mawhney,S, McLeod,M,P, McNeill,T,Z, Meenen,E,
Milosavljevic,A, Miner,G, Minja,E, Montemayor,J, Moore,S,
Morgan,M, Morris,K, Morris,S, Munidasa,M, Murphy,M, Nair,L,
Nankervis,C, Neal,D, Newton,N, Nguyen,N, Norris,S,
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Pasternak,S, Paul,H, Perez,A, Perez,L, Pfannkuch,C,
Plommer,F, Poindexter,A, Popovic,D, Primus,E, Pu,L, L,
Puzo,M, Quiroz,J, Rachlin,E, Reeves,K, Regier,M,A, Reigh,R,
Reilly,B, Reilly,M, Ren,Y, Reuter,M, Richards,S, Riggs,F,
Rives,C, Rodkey,T, Rojas,A, Rose,M, Rose,R, Ruiz,S,J,
Sanders,W, Savary,G, Scherer,S, Scott,G, Shatsman,S, Shen,H,
Shetty,J, Shvartsbeyn,A, Sisson,I, Sitter,C,D, Smajs,D,
Sneed,A, Sodergren,E, Song,X,Z, Sorelle,R, Sosa,J,
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Taylor,T, Thomas,N, Thomas,S, Tingey,A, Trejos,Z, Usmani,K,
Valas,R, Vera,V, Villanana,D, Waldron,L, Walker,B, Wang,J,
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Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von
Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs,R.A.

TITLE

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Best Local Similarity 95.2%; Pred. No. 63;

Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 34510 CAGTGACATGCAGGTCTATCT 34530

RESULT 2

AC133226/c

LOCUS AC133226 258823 bp DNA linear HTG 15-NOV-2002

DEFINITION Rattus norvegicus clone CH230-329C22, *** SEQUENCING IN PROGRESS

ACCESSION AC133226

VERSION AC133226.3 GI:25007420

KEYWORDS HTG; HTGS PHASE2; HTGS DRAFT; HTGS_ENRICHED.

SOURCE Rattus norvegicus (Norway rat)

ORGANISM

Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

1 (bases 1 to 258823)

REFERENCE

Muzny, D., Marlet, Metzker, M., Lee, S., Abramson, S., Adams, C., Alder, J., Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D., Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F., Biewald, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Evans, C., Evans, C.A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., George, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamil, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hogues, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpach, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, U., Lorensheva, L., Louleghed, H., Lozado, R.J., Lu, X., Ma, J., Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhney, S., McLeod, W.P., McNeill, T.Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwaokemele, O., Okwuonu, G., Olarnpsagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C., Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L., Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J., Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartbeyn, A., Sisson, I., Sitter, C.D., Smajls, D., Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J., Steinle, M., Strong, R., Sutton, A., Svatek, A., Tabors, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tinney, A., Trejos, Z., Usmani, K., Valae, R., Vera, V., Villaseana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,

Williams, G., Willson, R., Wlarczyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, S., Zhao, S., Dunn, D., Von Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A.
Direct Submission
Unpublished
2 (bases 1 to 258823)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (08-SEP-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 258823)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (15-NOV-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Nov 15, 2002 this sequence version replaced gi:22771302.
The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: KGNW
Center clone name: CH230-329C22
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 187552 bases at least Q40
Consensus quality: 190690 bases at least Q30
Consensus quality: 192289 bases at least Q20
Estimated insert size: 192259; sum-of-contigs estimation
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 1 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
* 1 258823: contig of 258823 bp in length.
* Location/Qualifiers
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1 .1057
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site:

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misc_feature
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ORIGIN
Query Match 92.4%; Score 19.4; DB 2; Length 258823;
Best Local Similarity 95.2%; Pred.No. 62;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTTAGCT 21
|||||
Db 21404 CAGTGACATGCAGGCTTAGCT 21384

RESULT 3
CR352267/c
LOCUS
DEFINITION
CR352267 140394 bp DNA linear HTG 12-MAR-2004
Danio rerio clone DKEY-174N5, WORKING DRAFT SEQUENCE, 9 unordered
pieces
ACCESSION
CR352267 GI:145433391
VERSION
HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
KEYWORDS
Danio rerio (zebrafish)
SOURCE
Danio rerio
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
Cypriniformes; Cyprinidae; Danio.
1 (bases 1 to 140394)
McLay,K.
Direct Submission
Submitted (10-MAR-2004) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
zfsh-help@sanger.ac.uk Clone request: clonerequest@sanger.ac.uk
On Mar 13, 2004 this sequence version replaced gi:45381849.
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: zfsh-help@sanger.ac.uk
----- Project Information
Center project name: zK174N5
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Chemistry: dye-terminator; 100% of reads
Consensus quality: 138030 bases at least Q40
Consensus quality: 138659 bases at least Q30
Consensus quality: 139080 bases at least Q20
Insert size: 139594; sum-of-contigs
Insert size: 156923; 6.3% error; agarose-fp
Quality coverage: 10.76x in Q20 bases; sum-of-contigs Quality
coverage: 10.22x in Q20 bases; agarose-fp
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 9 contigs. The true order of the pieces
* is not known and their order in this sequence record is

```

```

* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* this record will be updated with the finished sequence.
* as soon as it is available and the accession number will
* be preserved.
* 1 4851: contig of 4851 bp in length
* 4852: gap of 100 bp
* 15778: contig of 10827 bp in length
* 15779: gap of 100 bp
* 15879: contig of 22290 bp in length
* 38169: gap of 100 bp
* 38269: contig of 16312 bp in length
* 54581: gap of 100 bp
* 54681: contig of 33256 bp in length
* 87937: gap of 100 bp
* 88037: contig of 6415 bp in length
* 94552: gap of 100 bp
* 94552: contig of 12961 bp in length
* 107513: gap of 100 bp
* 107513: contig of 5237 bp in length
* 107613: gap of 100 bp
* 112850: contig of 27445 bp in length.
* 112950: Location/Qualifiers
1..140394
/organism="Danio rerio"
/mol_type="genomic DNA"
/db_xref="taxon:7955"
/clone="DKEY-174N5"
/clone_lib="DanioKey"
1..4851
/notes="assembly fragment:00165
fragment_chain:1"
4952..15778
/notes="assembly fragment:00354
fragment_chain:1"
15879..38168
/notes="assembly fragment:01046
fragment_chain:1"
38269..54580
/notes="assembly fragment:00715
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54681..87936
/notes="assembly fragment:01919
fragment_chain:1"
88037..94451
/notes="assembly fragment:00241
fragment_chain:1"
94552..107512
/notes="assembly fragment:00500
fragment_chain:1"
107613..112849
/notes="assembly fragment:00118
fragment_chain:2"
112950..140394
/notes="assembly fragment:01411
fragment_chain:2"

ORIGIN
Query Match 87.6%; Score 18.4; DB 2; Length 140394;
Best Local Similarity 95.0%; Pred.No. 2.1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 AGTGACATGCAGGCTTAGCT 21
|||||
Db 74881 AGTGACATGCAGGCTTAGCT 74862

RESULT 4
AP000772/c
LOCUS
DEFINITION
Homo sapiens chromosome 11 clone CMB9-7B14 map 11q22, WORKING DRAFT
SEQUENCE, 27 unordered pieces.
ACCESSION
AP000772

```


VERSION
KEYWORDS
SOURCE
ORGANISM

AP000772.2 GI:8118931
HTG; HTGS_PHASE1; HTGS_DRAFT.
Homo sapiens (human)
Homo sapiens
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 155858)

REFERENCE
AUTHORS

Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.

TITLE
JOURNAL

Homo sapiens 155,858 genomic DNA of 11q22
Published Only in DataBase (1999)

REFERENCE
AUTHORS

2 (bases 1 to 155858)
Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.

TITLE
JOURNAL

Submitted (25-NOV-1999) Masahira Hattori, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC),
Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555,
Japan (E-mail:hattori@gsc.riken.go.jp,
URL:http://hgp.gsc.riken.go.jp/, Tel:81-42-778-9923,
Fax:81-42-778-9924)
On May 31, 2000 this sequence version replaced gi:6997610.

COMMENT

----- Genome Center
Center: RIKEN Genomic Sciences Center (GSC)
Center code: RIKEN
Web site: http://hgp.gsc.riken.go.jp/
Contact: hattori@gsc.riken.go.jp
----- Project Information
Center project name: HumDraft11
Center clone name: CMB9-7B14

----- Summary Statistics
Sequencing vector: PCR products; 100% of reads
Chemistry: Dye-terminator ET-amerham; 100% of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 135660 bases at least Q40
Consensus quality: 145354 bases at least Q30
Consensus quality: 150658 bases at least Q20
Insert size: 153258; sum-of-contigs
Quality coverage: 4.31x in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently consists of
27 contigs. The true order of the pieces is not known and their
order in this sequence record is arbitrary. Gaps between the
contigs are represented as runs N, but the exact sizes of the gaps
are unknown. This record will be updated with the finished sequence
as soon as it is available and the accession number will be
preserved

1 18489 contig of 18489 bp in length
18590 36142 contig of 17553 bp in length
36243 49897 contig of 13655 bp in length
49998 63723 contig of 13726 bp in length
63824 76791 contig of 12968 bp in length
76892 85927 contig of 9036 bp in length
86028 93056 contig of 7029 bp in length
93157 99901 contig of 6745 bp in length
100002 106585 contig of 6584 bp in length
106686 111187 contig of 4502 bp in length
111288 115206 contig of 3919 bp in length
115307 119877 contig of 4571 bp in length
119978 124080 contig of 4103 bp in length
124181 127015 contig of 2835 bp in length
127116 129487 contig of 2372 bp in length
129588 133023 contig of 3436 bp in length
133124 136332 contig of 3209 bp in length
136433 139584 contig of 3252 bp in length
139685 142917 contig of 3132 bp in length
142917 143016 contig of 100 bp
143017 145169 contig of 2153 bp in length
145170 145269 contig of 100 bp
145270 147638 contig of 2369 bp in length
147639 147738 contig of 100 bp
147739 148903 contig of 1165 bp in length
148904 149003 contig of 100 bp
149004 150381 contig of 1378 bp in length
150382 150481 contig of 100 bp
150482 151677 contig of 1196 bp in length
151678 151777 contig of 100 bp
151778 153587 contig of 1810 bp in length
153588 153687 contig of 100 bp
153688 154698 contig of 1011 bp in length
154699 154798 contig of 100 bp
154799 155858 contig of 1060 bp in length.

FEATURES
source

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/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="11"
/map="11q22"
/clones="CMB9-7B14"
1..18489
/note="assembly_fragment"

misc_feature

1..18489
/note="assembly_fragment"

154799 155858 contig of 1060 bp in length
Sequence updated (26-May-2000).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 27 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 18489: contig of 18489 bp in length
* 18490 18589: gap of 100 bp
* 18590 36142: contig of 17553 bp in length
* 36143 36242: gap of 100 bp
* 36243 49897: contig of 13655 bp in length
* 49898 49997: gap of 100 bp
* 49998 63723: contig of 13726 bp in length
* 63724 63823: gap of 100 bp
* 63824 76791: contig of 12968 bp in length
* 76792 76891: gap of 100 bp
* 76892 85927: contig of 9036 bp in length
* 85928 86027: gap of 100 bp
* 86028 93056: contig of 7029 bp in length
* 93057 93157: gap of 100 bp
* 93157 99901: contig of 6745 bp in length
* 99902 100001: gap of 100 bp
* 100002 106585: contig of 6584 bp in length
* 106586 106685: gap of 100 bp
* 106686 111187: contig of 4502 bp in length
* 111188 111287: gap of 100 bp
* 111288 115206: contig of 3919 bp in length
* 115207 115306: gap of 100 bp
* 115307 119877: contig of 4571 bp in length
* 119878 119977: gap of 100 bp
* 119978 124080: contig of 4103 bp in length
* 124081 124180: gap of 100 bp
* 124181 127015: contig of 2835 bp in length
* 127016 127115: gap of 100 bp
* 127116 129487: contig of 2372 bp in length
* 129488 129587: gap of 100 bp
* 129588 133023: contig of 3436 bp in length
* 133024 133123: gap of 100 bp
* 133124 136332: contig of 3209 bp in length
* 136333 136432: gap of 100 bp
* 136433 139584: contig of 3252 bp in length
* 139585 139784: gap of 100 bp
* 139785 142916: contig of 3132 bp in length
* 142917 143016: gap of 100 bp
* 143017 145169: contig of 2153 bp in length
* 145170 145269: gap of 100 bp
* 145270 147638: contig of 2369 bp in length
* 147639 147738: gap of 100 bp
* 147739 148903: contig of 1165 bp in length
* 148904 149003: gap of 100 bp
* 149004 150381: contig of 1378 bp in length
* 150382 150481: gap of 100 bp
* 150482 151677: contig of 1196 bp in length
* 151678 151777: gap of 100 bp
* 151778 153587: contig of 1810 bp in length
* 153588 153687: gap of 100 bp
* 153688 154698: contig of 1011 bp in length
* 154699 154798: gap of 100 bp
* 154799 155858: contig of 1060 bp in length.


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fragment_chain:2"
misc_feature 100245..110317
/note="assembly_fragment:00215
fragment_chain:3"
misc_feature 110418..122844
/note="assembly_fragment:00326
fragment_chain:3"
misc_feature 122945..161516
/note="assembly_fragment:00891.0"
ORIGIN
Query Match 87.6%; Score 18.4; DB 2; Length 161516;
Best Local Similarity 95.0%; Pred. No. 2.1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 AGTGACATGCAGGCTAGCT 21
||||| ||||| ||||| ||||| |||||
Db 141651 AGTGACCTGCAGGCTAGCT 141632

RESULT 6
U21308
LOCUS U21308 40392 bp DNA linear INV 29-SEP-2004
DEFINITION Caenorhabditis elegans cosmid ZK1290, complete sequence.
ACCESSION U21308
VERSION U21308.1 GI:687795
KEYWORDS HTG.
SOURCE Caenorhabditis elegans
ORGANISM Caenorhabditis elegans
Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida;
Rhabditoides; Rhabditidae; Peloderinae; Caenorhabditis.
1 (bases 1 to 40392)
.
WormBase Consortium
Genome sequence of the nematode C. elegans: a platform for
investigating biology. The C. elegans Sequencing Consortium
Science 282 (5396), 2012-2018 (1998)
99069613
PUBMED 9851916
REFERENCE 2 (bases 1 to 40392)
AUTHORS Taich,A.
TITLE The sequence of C. elegans cosmid ZK1290
JOURNAL Unpublished (2001)
REFERENCE 3 (bases 1 to 40392)
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (13-JUL-1995) Department of Genetics, Washington
University, Genome Sequencing Center, 4444 Forest Park Avenue, St.
Louis, MO 63110, USA
4 (bases 1 to 40392)
REFERENCE 4 (bases 1 to 40392)
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (30-DEC-1997) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
5 (bases 1 to 40392)
REFERENCE 5 (bases 1 to 40392)
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (07-SEP-2001) Department of Genetics, Washington
University, Genome Sequencing Center, 4444 Forest Park Avenue, St.
Louis, MO 63110, USA
6 (bases 1 to 40392)
REFERENCE 6 (bases 1 to 40392)
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (29-MAY-2002) Department of Genetics, Washington
University, Genome Sequencing Center, 4444 Forest Park Avenue, St.
Louis, MO 63110, USA
7 (bases 1 to 40392)
REFERENCE 7 (bases 1 to 40392)
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (29-Oct-2002) Department of Genetics, Washington
University, Genome Sequencing Center, 4444 Forest Park Avenue, St.
Louis, MO 63110, USA

```

```

REFERENCE
AUTHORS
TITLE
JOURNAL
8 (bases 1 to 40392)
Waterston,R.
Direct Submission
Submitted (13-NOV-2002) Department of Genetics, Washington
University, Genome Sequencing Center, 4444 Forest Park Avenue, St.
Louis, MO 63110, USA
9 (bases 1 to 40392)
Waterston,R.
Direct Submission
Submitted (26-DEC-2002) Department of Genetics, Washington
University, Genome Sequencing Center, 4444 Forest Park Avenue, St.
Louis, MO 63110, USA
10 (bases 1 to 40392)
Waterston,R.
Direct Submission
Submitted (10-FEB-2003) Department of Genetics, Washington
University, Genome Sequencing Center, 4444 Forest Park Avenue, St.
Louis, MO 63110, USA
11 (bases 1 to 40392)
Waterston,R.
Direct Submission
Submitted (07-APR-2003) Department of Genetics, Washington
University, Genome Sequencing Center, 4444 Forest Park Avenue, St.
Louis, MO 63110, USA
12 (bases 1 to 40392)
Wilson,R.
Direct Submission
Submitted (05-NOV-2003) Department of Genetics, Washington
University, Genome Sequencing Center, 4444 Forest Park Avenue, St.
Louis, MO 63110, USA
13 (bases 1 to 40392)
Wilson,R.
Direct Submission
Submitted (12-DEC-2003) Department of Genetics, Washington
University, Genome Sequencing Center, 4444 Forest Park Avenue, St.
Louis, MO 63110, USA
14 (bases 1 to 40392)
Wilson,R.
Direct Submission
Submitted (06-AUG-2004) Department of Genetics, Washington
University, Genome Sequencing Center, 4444 Forest Park Avenue, St.
Louis, MO 63110, USA
15 (bases 1 to 40392)
WormBase Consortium
Direct Submission
Submitted (29-SEP-2004) Department of Genetics, Washington
University, Genome Sequencing Center, 4444 Forest Park Avenue, St.
Louis, MO 63110, USA
Submitted by:
Genome Sequencing Center
Department of Genetics, Washington University
St. Louis, MO 63110, USA, and
Sanger Centre, Hinxton Hall
Cambridge CB10 1RQ, England
email: submissions@watson.wustl.edu and jessesanger.ac.uk

```

NOTICE: This sequence may not be the entire insert of this clone. It may be shorter because we only sequence overlapping sections once, or longer because we provide a small overlap between neighboring submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one m13 subclone.

For a graphical representation of this clone sequence and its analysis see:

<http://www.wormbase.org/db/seq/sequence?name=ZK1290;class=Sequence>

* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.

1 949: contig of 949 bp in length
* 950 1049: gap of 100 bp
* 1050 2006: contig of 957 bp in length
* 2007 2106: gap of 100 bp
* 2107 3119: contig of 1013 bp in length
* 3120 3219: gap of 100 bp
* 3220 4172: contig of 953 bp in length
* 4173 4272: gap of 100 bp
* 4273 5244: contig of 972 bp in length
* 5245 5344: gap of 100 bp
* 5345 6288: contig of 944 bp in length
* 6289 6388: gap of 100 bp
* 6389 7363: contig of 975 bp in length
* 7364 7463: gap of 100 bp
* 7464 8447: contig of 984 bp in length
* 8448 8547: gap of 100 bp
* 8548 9561: contig of 1014 bp in length
* 9562 9661: gap of 100 bp
* 9662 10597: contig of 936 bp in length
* 10598 10697: gap of 100 bp
* 10698 11718: contig of 1021 bp in length
* 11719 11818: gap of 100 bp
* 11819 12831: contig of 1013 bp in length
* 12832 12931: gap of 100 bp
* 12933 13900: contig of 969 bp in length
* 13901 14000: gap of 100 bp
* 14001 14953: contig of 953 bp in length
* 14954 15053: gap of 100 bp
* 15054 16082: contig of 1029 bp in length
* 16083 16182: gap of 100 bp
* 16183 17195: contig of 1013 bp in length
* 17196 17295: gap of 100 bp
* 17296 18344: contig of 1049 bp in length
* 18345 18444: gap of 100 bp
* 18445 19385: contig of 941 bp in length
* 19386 19485: gap of 100 bp
* 19486 20455: contig of 970 bp in length
* 20456 20555: gap of 100 bp
* 20556 21548: contig of 993 bp in length
* 21549 21648: gap of 100 bp
* 21649 22631: contig of 983 bp in length
* 22632 22731: gap of 100 bp
* 22732 23705: contig of 974 bp in length
* 23706 23805: gap of 100 bp
* 23806 24762: contig of 957 bp in length
* 24763 24862: gap of 100 bp
* 24863 25835: contig of 973 bp in length
* 25836 25935: gap of 100 bp
* 25936 26918: contig of 983 bp in length
* 26919 27018: gap of 100 bp
* 27019 28003: contig of 985 bp in length
* 28004 28103: gap of 100 bp
* 28104 29109: contig of 1006 bp in length
* 29110 29209: gap of 100 bp
* 29210 30234: contig of 1025 bp in length
* 30235 30334: gap of 100 bp
* 30335 31327: contig of 993 bp in length
* 31328 31427: gap of 100 bp
* 31428 32386: contig of 971 bp in length
* 32389 32498: gap of 100 bp
* 32499 33501: contig of 1003 bp in length
* 33502 33601: gap of 100 bp
* 33602 34612: contig of 1011 bp in length
* 34613 34712: gap of 100 bp
* 34713 35718: contig of 1006 bp in length
* 35719 35818: gap of 100 bp
* 35819 36820: contig of 1002 bp in length
* 36821 36920: gap of 100 bp

36921 37914: contig of 994 bp in length
* 37915 38014: gap of 100 bp
* 38015 38989: contig of 975 bp in length
* 38990 39089: gap of 100 bp
* 39090 40045: contig of 956 bp in length
* 40046 40145: gap of 100 bp
* 40146 41107: contig of 962 bp in length
* 41108 41207: gap of 100 bp
* 41208 42219: contig of 1012 bp in length
* 42220 42319: gap of 100 bp
* 42320 43325: contig of 1006 bp in length
* 43326 43425: gap of 100 bp
* 43426 44439: contig of 1014 bp in length
* 44440 44539: gap of 100 bp
* 44540 45494: contig of 955 bp in length
* 45495 45594: gap of 100 bp
* 45595 46580: contig of 986 bp in length
* 46581 46680: gap of 100 bp
* 46681 47695: contig of 1015 bp in length
* 47696 47795: gap of 100 bp
* 47796 48789: contig of 994 bp in length
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* 48890 49831: contig of 942 bp in length
* 49832 49931: gap of 100 bp
* 49932 50926: contig of 995 bp in length
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* 52017 52116: gap of 100 bp
* 52117 53074: contig of 958 bp in length
* 53075 53174: gap of 100 bp
* 53175 54218: contig of 1044 bp in length
* 54219 54318: gap of 100 bp
* 54319 55335: contig of 1017 bp in length
* 55336 55435: gap of 100 bp
* 55436 56369: contig of 933 bp in length
* 56369 56468: gap of 100 bp
* 56469 57441: contig of 973 bp in length
* 57442 57541: gap of 100 bp
* 57542 58519: contig of 978 bp in length
* 58520 58619: gap of 100 bp
* 58620 59585: contig of 966 bp in length
* 59586 59685: gap of 100 bp
* 59686 60652: contig of 967 bp in length
* 60653 60752: gap of 100 bp
* 60753 61736: contig of 984 bp in length
* 61737 61836: gap of 100 bp
* 61837 62768: contig of 932 bp in length
* 62769 62868: gap of 100 bp
* 62869 63880: contig of 1012 bp in length
* 63881 63980: gap of 100 bp
* 63981 64974: contig of 994 bp in length
* 64975 65074: gap of 100 bp
* 65075 66085: contig of 1011 bp in length
* 66086 66185: gap of 100 bp
* 66186 67121: contig of 936 bp in length
* 67122 67221: gap of 100 bp
* 67222 68235: contig of 1014 bp in length
* 68236 68335: gap of 100 bp
* 68336 69321: contig of 986 bp in length
* 69322 69421: gap of 100 bp
* 69422 70456: contig of 1035 bp in length
* 70457 70556: gap of 100 bp
* 70557 71570: contig of 1014 bp in length
* 71571 71670: gap of 100 bp
* 71671 72655: contig of 985 bp in length.

FEATURES
source

1..72655
/organism="Mus musculus"
/mol_type="genomic DNA"
/db_xref="taxon:10090"

Query Match 84.8%; Score 17.8; DB 2; Length 72655;
Best Local Similarity 90.5%; Pred. No. 4.5e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCTAGCT 21
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 Db 26318 CTGTGACATGCAGATCTAGCT 26298

RESULT 8

AL831718/c
 LOCUS
 DEFINITION
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM

AL831718 116076 bp DNA linear ROD 15-NOV-2002
 Mouse DNA sequence from clone RP23-146020 on chromosome X, complete
 sequence.
 AL831718
 AL831718.6 GI:25137019
 HTG.
 Mus musculus (house mouse)
 Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 116076)
 Clark, S.

REFERENCE
 AUTHORS
 TITLE
 JOURNAL

Direct Submission
 Submitted (13-AUG-2002) Wellcome Trust Sanger Institute, Hinxton,
 Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
 humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
 On Nov 19, 2002 this sequence version replaced gi:22213735.
 ----- Genome Center
 Center: Wellcome Trust Sanger Institute
 Center code: SC
 Web site: http://www.sanger.ac.uk
 Contact: humquery@sanger.ac.uk

COMMENT

During sequence assembly data is compared from overlapping clones.
 Where differences are found these are annotated as variations
 together with a note of the overlapping clone name. Note that the
 variation annotation may not be found in the sequence submission
 corresponding to the overlapping clone, as we submit sequences with
 only a small overlap as described above.
 This sequence was finished as follows unless otherwise noted: all
 regions were either double-stranded or sequenced with an alternate
 chemistry or covered by high quality data (i.e., phred quality >=
 30); an attempt was made to resolve all sequencing problems, such
 as compressions and repeats; all regions were covered by at least
 one plasmid subclone or more than one M13 subclone; and the
 assembly was confirmed by restriction digest. The following
 abbreviations are used to associate primary accession numbers given
 in the feature table with their source databases: Em., EMBL; Sw.,
 SWISSPROT; Tr., TrEMBL; Wp., WORMPEP; Information on the WORMPEP
 database can be found at
 http://www.sanger.ac.uk/Projects/C_elegans/wormpep RP23-146020 is
 from the RPCI-23 Mouse PAC library
 constructed by the group of Pieter de Jong.
 For further details see http://www.chori.org/bacpac/home.htm
 VECTOR: pBACe3.6.

FEATURES
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 1. .116076
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 /mol_type="genomic DNA"
 /db_xref="taxon:10090"
 /chromosome="X"
 /clone="RP23-146020"
 /clone_lib="RPCI-23"

ORIGIN
 Query Match 84.8%; Score 17.8; DB 10; Length 116076;
 Best Local Similarity 90.5%; Pred. No. 4.2e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCTAGCT 21
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 Db 107376 CTGTGACATGCAGATCTAGCT 107356

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCTAGCT 21
 | ||||| ||||| |||||
 Db 107376 CTGTGACATGCAGATCTAGCT 107356

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

RESULT 9

AC148761/c
 LOCUS
 DEFINITION

AC148761
 AC148761.1 GI:46063628
 HTG; HTGS_PHASE1; HTGS_ACTIVEPIN.
 Medicago truncatula (barrel medic)
 Medicago truncatula
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 rosids; eurosids I; Fabales; Fabaceae; Papilionoideae; Trifolieae;
 Medicago.
 1 (bases 1 to 125354)
 Town, C.D., Tallon, L.J., Arbogast, T., Althoff, R., Hine, E.,
 Monaghan, E., Smith, S.A., Utterback, T., Feldblyum, T. and Koo, H.
 Medicago truncatula BAC genomic sequence
 Unpublished
 2 (bases 1 to 125354)
 Town, C.D.

REFERENCE
 AUTHORS
 TITLE
 JOURNAL

Direct Submission
 Submitted (02-APR-2004) The Institute for Genomic Research, 9712
 Medical Center Dr, Rockville, MD 20850, USA

COMMENT

* NOTE: This is a 'working draft' sequence. It currently
 * consists of 14 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

1 1075: contig of 1075 bp in length
 1076 1175: gap of unknown length
 1176 2410: contig of 1235 bp in length
 2411 2510: gap of unknown length
 2511 29548: contig of 27038 bp in length
 29549 29648: gap of unknown length
 29649 32649: contig of 2721 bp in length
 32650 32659: gap of unknown length
 32660 32669: gap of unknown length
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 3

The RPCI-24 BAC library has been constructed by Pieter de Jong and coworkers (<http://www.chori.org>) from male C57BL/6J mouse spleen and/or brain genomic DNA. The clone and detailed information can be obtained from Pieter de Jong and coworkers at <http://www.chori.org>

NEIGHBORING SEQUENCE INFORMATION:
This sequence is the entire insert of the clone.

FEATURES
source
1. .139884
/organism="Mus musculus"
/mol_type="genomic DNA"
/db_xref="taxon:10090"
/map="15"
/chromosome="15"
/clone="RP24-193N24"
/clone_lib="RPCI-24"
1472..1532
/rpt_family="Alu"
repeat_region
2336..2478
/rpt_family="B4"
repeat_region
3625..3939
/rpt_family="MaLR"
repeat_region
4251..4445
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4623..4984
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6912..7089
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7105..7222
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7264..7331
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7640..7961
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9097..9389
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11701..11779
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13535..13735
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21199..21262
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21210..21311
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22533..22620

Db 42736 CAGTGACATGCAGGCTTACCT 42716
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RESULT 10
AC110379 139884 bp DNA linear ROD 05-NOV-2003
LOCUS Mus musculus BAC clone RP24-193N24 from 15, complete sequence.
AC110379
AC110379.3 GI:21536156
HTG.
Mus musculus (house mouse)
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 139884)
Haakenson, W. and Doebber, A.
The sequence of Mus musculus BAC clone RP24-193N24
Unpublished (2001)
2 (bases 1 to 139884)
WILSON, R.
Sequencing of Mus musculus
Unpublished (2001)
3 (bases 1 to 139884)
McPherson, J.D. and Waterston, R.H.
Direct Submission
Submitted (11-FEB-2002) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
4 (bases 1 to 139884)
McPherson, J.D. and Waterston, R.H.
Direct Submission
Submitted (08-APR-2002) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
5 (bases 1 to 139884)
McPherson, J.D. and Waterston, R.H.
Direct Submission
Submitted (21-JUN-2002) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
6 (bases 1 to 139884)
Wilson, R.
Direct Submission
Submitted (05-NOV-2003) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Jun 21, 2002 this sequence version replaced gi:20069820.
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu>
Contact: submissions@wustl.edu
----- Summary Statistics

Center project name: M_BB0193N24

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. Wes Warren, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu>

SOURCE INFORMATION:

/rpt family="B4"
23026. .23125
/rpt family="Alu"
23974. .24135
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24176. .24297
/rpt family="Alu"
24344. .24414
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25063. .25207
/rpt family="MaLR"
25971. .27396
/rpt family="MaLR"
27790. .28076
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28465. .28682
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30110. .30243
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32501. .32551
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34609. .34755
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/rpt family="B4"
36028. .36213
/rpt family="B2"
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/rpt family="B4"
38777. .38983
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41364. .41539
/rpt family="B2"
42540. .42702
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/rpt family="B4"
43887. .44000
/rpt family="B4"
44624. .44961
/rpt family="MaLR"
45125. .45221
/rpt family="MER2_type"
46183. .46321

Query Match 84.8%; Score 17.8; DB 10; Length 139884;
Best Local Similarity 90.3%; Pred. No. 4.1e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 CAGTGACATGCAGGCTAGCT 21
||||| ||||| ||||| |||||
Db 9945 CAGTGACTGCAGGGCTAGCT 9965
RESULT 11
AC102916/c
LOCUS AC102916 linear 144142 bp DNA ROD 29-SEP-2004

DEFINITION
AC102916
VERSION AC102916.5 GI:52839774
KEYWORDS HTG.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
Birren,B., Nussbaum,C. and Lander,E.
1 (bases 1 to 144142)
Mus musculus chromosome 5, clone RP24-274118
Unpublished
2 (bases 1 to 144142)
Birren,B., Linton,L., Nussbaum,C., Lander,E., Ali,A., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Boguslavskiy,L., Boukhgalter,B.,
Brown,A., Camarata,J., Campopiano,A., Chang,J.J., Chazaro,B.,
Choepel,Y., Collangelo,M., Collins,S., Collymore,A., Cook,A.,
Cooke,P., DeArellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,
Ferreira,P., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S.,
Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,
Jones,C., Kamat,A., Karatas,A., Kells,C., LaRocque,K.,
Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Liu,G.,
MacClean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C.,
McCarthy,M., McEwan,P., McKernan,K., McPheeters,R., Meldrim,J.,
Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C.,
Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neil,D.,
Olliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,
Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,
Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupback,R.,
Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

TITLE
JOURNAL
REFERENCE
AUTHORS

Direct Submission
Submitted (23-NOV-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 144142)
Birren,B., Nussbaum,C., Lander,E., Abouelleil,A., Allen,N.,
Anderson,M., Anderson,S., Arachchi,H.M., Barna,N., Bastien,V.,
Bloom,T., Boguslavskiy,L., Boukhgalter,B., Camarata,J., Chang,J.,
Choepel,Y., Collymore,A., Cook,A., Cooke,P., Corum,B.,
DeArellano,K., Diaz,J.S., Dodge,S., Dooley,K., Dorris,L.,
Erickson,J., Faro,S., Ferreira,P., FitzGerald,M., Gage,D.,
Galagan,J., Gardyna,S., Graham,L., Grand-Pierre,N., Hafez,N.,
Hagopian,D., Hagos,B., Hall,J., Horton,L., Hulme,W., Iliev,I.,
Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Landers,T.,
Levine,R., Lindblad-Toh,K., Liu,G., Liu,X., Lui,A., Mabbitt,R.,
MacClean,C., Macdonald,P., Major,J., Manning,J., Matthews,C.,
McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V.,
Murphy,T., Naylor,J., Nguyen,C., Nguyen,T., Nicol,R., Norbu,C.,
O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Rachupka,A., Ramasamy,U., Raymond,C.,
Retta,R., Rise,C., Rogov,P., Roman,J., Schauer,S., Schupback,R.,
Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N.,
Stojanovic,N., Stubbs,M., Talamas,J., Tesfaye,S., Theodore,J.,
Topham,K., Travers,M., Vassiliev,H., Venkataraman,V.S., Viel,R.,
Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L.,
Zimmer,A. and Zody,M.

TITLE
JOURNAL
REFERENCE
AUTHORS

Direct Submission
Submitted (21-AUG-2004) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
4 (bases 1 to 144142)
Birren,B., Nussbaum,C., Lander,E., Abouelleil,A., Allen,N.,
Anderson,M., Anderson,S., Arachchi,H.M., Barna,N., Bastien,V.,
Bloom,T., Boguslavskiy,L., Boukhgalter,B., Camarata,J., Chang,J.,
Choepel,Y., Collymore,A., Cook,A., Cooke,P., Corum,B.,
DeArellano,K., Diaz,J.S., Dodge,S., Dooley,K., Dorris,L.,
Erickson,J., Faro,S., Ferreira,P., FitzGerald,M., Gage,D.,
Galagan,J., Gardyna,S., Graham,L., Grand-Pierre,N., Hafez,N.,
Hagopian,D., Hagos,B., Hall,J., Horton,L., Hulme,W., Iliev,I.,
Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Landers,T.,


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REFERENCE
AUTHORS
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
1 (bases 1 to 162289)
Ahter,N., Antonellis,A., Ayele,K., Becketrom-Sternberg,S.M.,
Benjamin,B., Blakesley,R.W., Bouffard,G.G., Breen,K., Brinkley,C.,
Brooks,S., Dietrich,N.L., Granite,S., Guan,X., Gupta,J.,
Haghighi,P., Hansen,N., Ho,S.-L., Idol,J.R., Karlins,E., Laric,P.,
Lee-Lin,S.-Q., Legaspi,R., Maduro,Q.L., Maduro,V.B.,
Margulies,E.H., Masiello,C., Maskeri,B., Mastrrian,S.D.,
McCloskey,J.C., McDowell,J., Paguirigan,C., Pearson,R.,
Portnoy,M.E., Prasad,A., Schueler,M.G., Stantripop,S., Thomas,J.W.,
Thomas,P.J., Touchman,J.W., Tsurgeon,C., Vogt,J.L., Walker,M.A.,
Wetherby,K.D., Wiggins,L., Young,A., Zhang,L.-H. and Green,E.D.
NISC Comparative Sequencing Initiative
Unpublished
2 (bases 1 to 162289)
Green,E.D.
Direct Submission
Submitted (12-OCT-2001) NIH Intramural Sequencing Center, 8717
Grovemont Circle, Gaithersburg, MD 20877, USA
3 (bases 1 to 162289)
Green,E.D.
Direct Submission
Submitted (22-FEB-2002) NIH Intramural Sequencing Center, 8717
Grovemont Circle, Gaithersburg, MD 20877, USA
4 (bases 1 to 162289)
Green,E.D.
Direct Submission
Submitted (26-JUL-2002) NIH Intramural Sequencing Center, 8717
Grovemont Circle, Gaithersburg, MD 20877, USA
5 (bases 1 to 162289)
Green,E.D.
Direct Submission
Submitted (28-JAN-2003) NIH Intramural Sequencing Center, 8717
Grovemont Circle, Gaithersburg, MD 20877, USA
On Jul 26, 2002 this sequence version replaced gi:18860664.
----- Genome Center
Center: NIH Intramural Sequencing Center
Center code: NISC
Web site: http://www.nisc.nih.gov
Contact: nisc.zoo@nhgri.nih.gov
----- Project Information
Center project name: cnz
Center clone name: 254G01

This sequence was finished as follows unless otherwise noted:
all regions were double-stranded, sequenced with an
alternate chemistry, or covered by high quality data
(i.e., phred quality >= 30); an attempt was made to resolve
all sequencing problems, such as compressions and repeats;
all regions were covered by at least one plasmid subclone
or more than one M13 subclone; and the assembly was confirmed
by restriction digest.

CLONE LENGTH: This sequence represents the entire insert of
this clone unless otherwise noted. If there are overlapping
clones, the overlaps are noted in the beginning and end of
the Features section.
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Mammalia; Eutheria; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 162560)
Waterston,R.H.
The sequence of Homo sapiens clone
Unpublished
2 (bases 1 to 162560)
Waterston,R.H.
Direct Submission
Submitted (16-MAY-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Jun 30, 2000 this sequence version replaced gi:8469066.
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information
Center project name: H.NH0712115
----- Summary Statistics
Sequencing vector: M13; 100%
Sequencing vector: plasmid; 0%
Chemistry: Dye-primer ET; 100% of reads
Chemistry: Dye-terminator Big Dye; 0% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 145600 bases at least Q40
Consensus quality: 151065 bases at least Q30
Consensus quality: 153390 bases at least Q20
Insert size: 173000; agarose-fp
Insert size: 159360; sum-of-contigs
Quality coverage: 3.35 in Q20 bases; agarose-fp
Quality coverage: 3.73 in Q20 bases; sum-of-contigs
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* NOTE: This is a 'working draft' sequence. It currently
* consists of 33 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
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* 1 1483: contig of 1483 bp in length
* 1484 1583: gap of unknown length
* 1584 3194: contig of 1611 bp in length
* 3195 3294: gap of unknown length
* 3295 4473: contig of 1178 bp in length
* 4473 4572: gap of unknown length
* 4572 6120: contig of 1548 bp in length
* 6121 6221: gap of unknown length
* 6221 8046: contig of 1826 bp in length

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Lindblad-Toh, K., Liu, G., Maclean, C., Macdonald, P., Major, J., Matthews, C., McCarthy, M., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neil, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P., Roman, J., Roy, A., Schauer, S., Schuback, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J., Testaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission
Submitted (07-JAN-2003) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jan 7, 2003 this sequence version replaced gi:27476180.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L21648
Center clone name: 421_P_23

FEATURES

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			Gaps	0;

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DB	842	CAGTCACATGCAGGCTCAGCT	822

RESULT	15
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LOCUS	AC015468
DEFINITION	Homo sapiens chromosome 8, clone RP11-369E15, complete sequence.
ACCESSION	AC015468
VERSION	AC015468.5 GI:138999433

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KEYWORDS
SOURCE      Homo sapiens (human)
ORGANISM    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE
AUTHORS    Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
TITLE      1 (bases 1 to 189662)
JOURNAL    Birren,B., Linton,L., Nusbaum,C. and Lander,E.
REFERENCE  1 (bases 1 to 189662)
AUTHORS    Unpublished
            2 (bases 1 to 189662)
            Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
            Baldwin,J., Barna,N., Beckerly,R., Boguslavskiy,L., Boukhgalter,B.,
            Brown,A., Castle,A., Colangelo,M., Collins,S., Collymore,A.,
            Cooke,P., DeArelano,K., Dewar,K., Domino,M., Donelan,D., Doyle,M.,
            Ferreira,P., FitzHugh,W., Forrest,C., Funke,R., Gage,D.,
            Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
            Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
            Lehoczy,J., Lieu,C., Locke,K., Macdonald,P., Marquis,N.,
            McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrim,J.,
            Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
            Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severy,P.,
            Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
            Tesfaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,
            Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.

TITLE      Direct Submission
JOURNAL    Submitted (16-NOV-1999) Whitehead Institute/MIT Center for Genome
            Research, 320 Charles Street, Cambridge, MA 02141, USA
REFERENCE  3 (bases 1 to 189662)
AUTHORS    Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,
            Barna,N., Bastien,V., Boguslavskiy,L., Boukhgalter,B., Brown,A.,
            Camarata,J., Campopiano,A., Chang,J., Choepel,Y., Colangelo,M.,
            Collins,S., Collymore,A., Cooke,P., DeArelano,K., Dewar,K.,
            Diaz,J.S., Dodge,S., Faro,S., Ferreira,P., FitzHugh,W., Gage,D.,
            Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,
            Grand-Pierre,N., Hagos,B., Heaford,A., Horton,L., Hulme,W.,
            Iliev,I., Johnson,R., Jones,C., Karatas,A., LaRocque,K.,
            Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Liu,G.,
            McLean,C., Macdonald,P., Marquis,N., Matthews,C., McCarthy,M.,
            McEwan,P., McKernan,K., McPeeters,R., Meldrim,J., Meneus,L.,
            Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Norbu,C.,
            Norman,C.H., O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J.,
            Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C.,
            Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J.,
            Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupback,R., Seaman,S.,
            Severy,P., Sougnez,C., Spencer,B., Stange-Thomann,N.,
            Stojanovic,N., Straus,C., Subramanian,A., Talamas,J., Tesfaye,S.,
            Theodore,J., Travers,M., Travis,N., Triglio,J., Vassiliev,H.,
            Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
            Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

TITLE      Direct Submission
JOURNAL    Submitted (01-MAY-2001) Whitehead Institute/MIT Center for Genome
            Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT    On May 1, 2001 this sequence version RepeatMasker:
            All repeats were identified using RepeatMasker:
            Smith, A.F.A. & Green, P. (1996-1997)
            http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information
Center project name: L2466
Center clone name: 369_E_15

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   /map="8"
   /clone="RP11-369E15"

/clone_lib="RPC1-11 Human Male BAC"
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15167. .15332
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FEATURES
source

contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: KATP

Center clone name: CH230-304B3

----- Summary Statistics

Assembly program: Phrap; version 0.990329

Consensus quality: 191737 bases at least Q40

Consensus quality: 193479 bases at least Q30

Consensus quality: 194485 bases at least Q20

Estimated insert size: 194733; sum-of-contigs estimation

Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently
* consists of 5 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

* 1 191071: contig of 191071 bp in length
* 191072 191171: gap of unknown length
* 191172 192706: contig of 1535 bp in length
* 192707 192806: gap of unknown length
* 192807 194126: contig of 1320 bp in length
* 194127 194226: gap of unknown length
* 194227 196128: contig of 1902 bp in length
* 196129 196228: gap of unknown length
* 196229 197796: contig of 1568 bp in length.

FEATURES

source

1. 197796

/organism="Rattus norvegicus"

/mol_type="genomic DNA"

/db_xref="taxon:10116"

/clone="CH230-304B3"

1. 1063

/notes="wgs end_extension"

clone_end:T7"

1565. 2730

/notes="wgs end_extension"

clone_end:T7"

3873. 4716

/notes="clone_boundary"

clone_end:T7"

site:

end_sequence:BZ204887"

ORIGIN

Query Match 84.8%; Score 17.8; DB 2; Length 197796;

Best Local Similarity 90.5%; Pred. No. 4e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CACTGACATGCAGGCTTAGCT 21

Db 192834 CTGTGATATGCAGGCTTAGCT 192854

RESULT 17

AC129792/c

LOCUS

AC129792

Rattus norvegicus

***, 5 unordered pieces.

AC129792

DEFINITION

***, 5 unordered pieces.

AC129792

197796 bp DNA linear HTG 19-NOV-2002

Rattus norvegicus clone CH230-304B3, *** SEQUENCING IN PROGRESS

***, 5 unordered pieces.

AC129792

VERSION

KEYWORDS

SOURCE

ORGANISM

AC129792.4 GI:25073629

HTG; HTGS PHASE1; HTGS DRAFT; HTGS_ENRICHED.

Rattus norvegicus (Norway rat)

Rattus norvegicus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

1 (bases 1 to 197796)

REFERENCE

AUTHORS

Muzny,D,Marie, Metzker,M, Lee, Abramson, S., Adams, C., Alder, J., Allen, C., Allen, H., Alebrooks, S., Amin, A., Anguiano, D., Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F., Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Crease, A., D'Souza, L., Davila, M., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Eugene, C., Evans, C. A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C. M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregeorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamil, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S. L., Hodgson, A., Hoques, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C. L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensuhewa, L., Loulseged, H., Lozado, R. J., Lu, X., Ma, J., Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhiney, S., McLeod, M. P., McNeill, T. Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwaokelemeh, O., Okwuonu, G., Olarnpunsagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C., Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L., L., Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M. A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S. J., Sanders, W., Savary, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C. D., Smajls, D., Sneed, A., Sodergren, E., Song, X. Z., Sorelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A., Svatek, A., Taber, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K., Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wiecezyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, S., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D. R., Holt, R. A., Smith, H. O., Weinstock, G. and Gibbs, R. A.

Direct Submission

Unpublished

2 (bases 1 to 197796)

Worley, K. C.

Direct Submission

Submitted (03-AUG-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 197796)

Rat Genome Sequencing Consortium.

Direct Submission

Submitted (19-NOV-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

On Nov 19, 2002 this sequence version replaced gi:23915295.

The sequence in this assembly is a combination of BAC based reads

and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: KATP
Center clone name: CH230-304B3
----- Summary Statistics

Assembly program: Phrap; version 0.990329
Consensus quality: 191737 bases at least Q40
Consensus quality: 193479 bases at least Q30
Consensus quality: 194485 bases at least Q20
Estimated insert size: 194733; sum-of-contigs estimation
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 5 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved

* 1 191071: contig of 191071 bp in length
* 191072 191171: gap of unknown length
* 191172 192706: contig of 1535 bp in length
* 192707 192806: gap of unknown length
* 192807 194126: contig of 1320 bp in length
* 194127 194226: gap of unknown length
* 194227 196128: contig of 1902 bp in length
* 196129 196228: gap of unknown length
* 196229 197796: contig of 1568 bp in length.

FEATURES

source
 1. .197796
 /organism="Rattus norvegicus"
 /mol_type="genomic DNA"
 /db_xref="taxon:10116"
 /clone="CH230-304B3"
misc_feature
 1. .1063
 /notes="wgs end_extension
 clone_end:T7"
misc_feature
 1565. .2730
 /notes="wgs end_extension
 clone_end:T7"
misc_feature
 3873. .4716
 /notes="clone boundary
 clone_end:T7"
 site:
 end_sequence:BZ204887"

ORIGIN

Query Match 84.8%; Score 17.8; DB 2; Length 197796;
Best Local Similarity 90.5%; Pred. No. 4e-02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Oy 1 CACTGACATGCAGGCTTAGCT 21
 |||||
Db 97483 CTGTGATATGCAGGCTTAGCT 97463

RESULT 18
AC087221
LOCUS

DEFINITION

AC087221

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Homo sapiens

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

AC087221 203690 bp DNA linear HTG 24-MAY-2002
Homo sapiens chromosome 8 clone RP11-712115 map 8, WORKING DRAFT
SEQUENCE, 34 ordered pieces.
AC087221
AC087221-2 GI:21166223
HTG; HTGS_PHASE2; HTGS_DRAFT; HTGS_FULLTOP.
Homo sapiens (human)

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 203690)

Birren,B., Linton,L., Nusbaum,C. and Lander,E.
Homo sapiens chromosome 8, clone RP11-712115
Unpublished
2 (bases 1 to 203690)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Boguslavskiy,L., Bouckhalter,B., Brown,A.,
Camarata,J., Campopiano,A., Choepel,Y., Colangelo,M., Collins,S.,
Collamore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S.,
Dodge,S., Faro,S., Ferreira,P., FitzHugh,W., Gage,D., Galagan,J.,
Gardyna,S., Ginde,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,
Jones,C., Karatas,A., LaRocque,K., Lamazares,R., Landers,T.,
Lehoczky,J., Levine,R., Liu,G., MacLean,C., Macdonald,P.,
Marquis,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K.,
McPheeters,R., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V.,
Murphy,T., Naylor,J., Nguyen,C., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Pollara,V., Raymond,C., Retta,R.,
Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J., Rosetti,M.,
Roy,A., Santos,R., Schauer,S., Schupback,R., Seaman,S., Severy,P.,
Sounges,C., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Travers,M., Travis,N., Trigilio,J., Vassiliev,H., Viel,R., Vo,A.,
Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.

Direct Submission
Submitted (16-DEC-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 203690)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L.,
Bouckhalter,B., Brown,A., Camarata,J., Campopiano,A., Chang,J.,
Chazaro,B., Choepel,Y., Colangelo,M., Collins,S., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., FitzGerald,M., FitzHugh,W., Gage,D.,
Galagan,J., Gardyna,S., Ginde,S., Gord,S., Goyette,M., Graham,L.,
Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I.,
Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Lakocque,K.,
Lamazares,R., Landers,T., Lehoczky,J., Levine,R., Lindblad-Toh,K.,
Liu,G., MacLean,C., Macdonald,P., Major,J., Marquis,N.,
Matthews,C., McCarthy,M., McEwan,P., McKernan,K., Meldrim,J.,
Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C.,
Nicoli,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P.,
O'Neil,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,
Pollara,V., Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C.,
Rogov,P., Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S.,
Schupback,R., Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N.,
Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S.,
Theodore,J., Topham,K., Travers,M., Travis,N., Trigilio,J.,
Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

Direct Submission
Submitted (24-MAY-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On May 24, 2002 this sequence version replaced gi:11875303.

All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: <http://www-seq.wi.mit.edu>

Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L11638

Center clone name: 712_1_15

----- Summary Statistics

Sequencing vector: Plasmid; n/a; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 190084 bases at least Q40

Consensus quality: 196705 bases at least Q30

Consensus quality: 199369 bases at least Q20

Insert size: 176000; agarose-fp

Insert size: 200390; sum-of-contigs

Quality coverage: 9.6 in Q20 bases; agarose-fp

Quality coverage: 8.5 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 34 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submittor.

* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.

* 1 460: contig of 460 bp in length

* 461 560: gap of 100 bp

* 561 1196: contig of 636 bp in length

* 1197 1296: gap of 100 bp

* 1297 2122: contig of 826 bp in length

* 2123 2222: gap of 100 bp

* 2223 2859: contig of 647 bp in length

* 2870 2969: gap of 100 bp

* 2970 3620: contig of 651 bp in length

* 3621 3720: gap of 100 bp

* 3721 4314: contig of 594 bp in length

* 4315 4414: gap of 100 bp

* 4415 5152: contig of 738 bp in length

* 5153 5252: gap of 100 bp

* 5253 6209: contig of 957 bp in length

* 6210 6309: gap of 100 bp

* 6310 7103: contig of 794 bp in length

* 7104 7203: gap of 100 bp

* 7204 7937: contig of 734 bp in length

* 7938 8037: gap of 100 bp

* 8038 9006: contig of 969 bp in length

* 9007 9107: gap of 100 bp

* 9107 9906: contig of 800 bp in length

* 9907 10006: gap of 100 bp

* 10007 11070: contig of 1064 bp in length

* 11071 11170: gap of 100 bp

* 11171 12297: contig of 1127 bp in length

* 12298 12397: gap of 100 bp

* 12398 13415: contig of 1018 bp in length

* 13416 13515: gap of 100 bp

* 13516 14231: contig of 716 bp in length

* 14232 14331: gap of 100 bp

* 14332 15596: contig of 1265 bp in length

* 15597 15696: gap of 100 bp

* 15697 16462: contig of 766 bp in length

* 16463 16562: gap of 100 bp

* 16563 17742: contig of 1180 bp in length

* 17743 17842: gap of 100 bp

* 17843 18857: contig of 1015 bp in length

* 18858 18957: gap of 100 bp

* 18958 20490: contig of 1533 bp in length

* 20491 20590: gap of 100 bp

* 20591 22210: contig of 1620 bp in length

* 22211 22310: gap of 100 bp

* 22311 23851: contig of 1541 bp in length

* 23852 23951: gap of 100 bp

* 23952 25684: contig of 1733 bp in length

* 25685 25784: gap of 100 bp

* 25785 27006: contig of 1222 bp in length

* 27007 27106: gap of 100 bp

* 27107 28616: contig of 1510 bp in length

* 28617 28716: gap of 100 bp

* 28717 30184: contig of 1468 bp in length

* 30185 30284: gap of 100 bp

* 30285 31303: contig of 1019 bp in length

* 31304 31403: gap of 100 bp

* 31404 32917: contig of 1514 bp in length

* 32918 33017: gap of 100 bp

* 33018 42030: contig of 9013 bp in length

* 42031 42130: gap of 100 bp

* 42131 49012: contig of 6882 bp in length

* 49013 49112: gap of 100 bp

* 49113 62240: contig of 13128 bp in length

* 62241 62340: gap of 100 bp

* 62341 86795: contig of 24455 bp in length

* 86796 86895: gap of 100 bp

* 86896 203690: contig of 116795 bp in length.

FEATURES

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/db_xref="taxon:9606"

/chromosome="8"

/map="8"

/clone="RP11-712115"

/clone_lib="RPC1-11 Human Male BAC"

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clone_end:SP6

vector_side:left"

561..1196

/note="assembly_fragment"

1297..2122

/note="assembly_fragment"

2223..2869

/note="assembly_fragment"

2970..3620

/note="assembly_fragment"

3721..4314

/note="assembly_fragment"

4415..5152

/note="assembly_fragment"

5253..6209

/note="assembly_fragment"

6310..7103

/note="assembly_fragment"

7204..7937

/note="assembly_fragment"

8038..9006

/note="assembly_fragment"

9107..9906

/note="assembly_fragment"

Query Match

Best Local Similarity 84.8%; Score 17.8; DB 2; Length 203690;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTCACATGCAGGCTCTAGCT 21

|||||

Db 190273 CAGTCACATGCAGGCTCTAGCT 190293

RESULT 19

AC115746

LOCUS

AC115746

DEFINITION

AC115746

ACCESSION

VERSION

AC115746 AC115746 214765 bp DNA linear ROD 29-JUL-2004

DEFINITION Mus musculus chromosome 15, clone RP23-3J8, complete sequence.

ACCESSION AC115746

VERSION AC115746.10 GI:50811761

HTG.	Mus musculus (house mouse)	
SOURCE	Mus musculus	
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.	
REFERENCE	1 (bases 1 to 214765)	
AUTHORS	Birren,B., Nusbaum,C. and Lander,E.	
TITLE	Mus musculus chromosome 15, clone RP23-3J8	
JOURNAL	Unpublished	
REFERENCE	2 (bases 1 to 214765)	
AUTHORS	Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B., Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B., Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S., Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., LaRocque,K., Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Lindblad-Toh,K., Liu,G., MacLean,C., Macdonald,P., Major,J., Marquis,N., Matthews,L., McCarthy,M., McEwan,T., Meldrim,J., Nelson,K., O'Neil,D., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., O'Connor,T., P., O'Donnell,P., Pierre,N., Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P., Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M., Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.	
TITLE	Direct Submission	
JOURNAL	Submitted (22-MAR-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA	
REFERENCE	3 (bases 1 to 214765)	
AUTHORS	Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N., Anderson,M., Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B., Camarata,J., Chang,J., Choepel,Y., Collymore,A., Cooke,A., Cooke,P., Corum,B., Dearellano,K., Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S., Graham,L., Grand-Pierre,N., Hafez,N., Hagopian,D., Hagos,B., Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K., Liu,G., Liu,X., Lui,A., Mabbitt,R., MacLean,C., Macdonald,P., Major,J., Manning,J., Matthews,C., McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nguyen,T., Nicol,R., Norbu,C., O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P., Roman,J., Schauer,S., Schupback,R., Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M., Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.	
TITLE	Direct Submission	
JOURNAL	Submitted (22-MAR-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA	
REFERENCE	4 (bases 1 to 214765)	
AUTHORS	Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N., Anderson,M., Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B., Camarata,J., Chang,J., Choepel,Y., Collymore,A., Cooke,A., Cooke,P., Corum,B., Dearellano,K., Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S., Graham,L., Grand-Pierre,N., Hafez,N., Hagopian,D., Hagos,B., Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K., Liu,G., Liu,X., Lui,A., Mabbitt,R., MacLean,C., Macdonald,P., Major,J., Manning,J., Matthews,C., McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nguyen,T., Nicol,R., Norbu,C., O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P., Roman,J., Schauer,S., Schupback,R., Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M., Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.	
TITLE	Direct Submission	
JOURNAL	Submitted (03-JUN-2004) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA	
REFERENCE	5 (bases 1 to 214765)	
AUTHORS	Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N., Anderson,M., Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B., Camarata,J., Chang,J., Choepel,Y., Collymore,A., Cooke,A., Cooke,P., Corum,B., Dearellano,K., Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S., Graham,L., Grand-Pierre,N., Hafez,N., Hagopian,D., Hagos,B., Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K., Liu,G., Liu,X., Lui,A., Mabbitt,R., MacLean,C., Macdonald,P., Major,J., Manning,J., Matthews,C., McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nguyen,T., Nicol,R., Norbu,C., O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P., Roman,J., Schauer,S., Schupback,R., Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M., Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.	
TITLE	Direct Submission	
JOURNAL	Submitted (03-JUN-2004) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA	
REFERENCE	6 (bases 1 to 214765)	
AUTHORS	Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N., Anderson,M., Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B., Camarata,J., Chang,J., Choepel,Y., Collymore,A., Cooke,A., Cooke,P., Corum,B., Dearellano,K., Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S., Graham,L., Grand-Pierre,N., Hafez,N., Hagopian,D., Hagos,B., Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K., Liu,G., Liu,X., Lui,A., Mabbitt,R., MacLean,C., Macdonald,P., Major,J., Manning,J., Matthews,C., McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nguyen,T., Nicol,R., Norbu,C., O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P., Roman,J., Schauer,S., Schupback,R., Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M., Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.	
TITLE	Direct Submission	
JOURNAL	Submitted (03-JUN-2004) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA	
REFERENCE	7 (bases 1 to 214765)	
AUTHORS	Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N., Anderson,M., Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B., Camarata,J., Chang,J., Choepel,Y., Collymore,A., Cooke,A., Cooke,P., Corum,B., Dearellano,K., Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S., Graham,L., Grand-Pierre,N., Hafez,N., Hagopian,D., Hagos,B., Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K., Liu,G., Liu,X., Lui,A., Mabbitt,R., MacLean,C., Macdonald,P., Major,J., Manning,J., Matthews,C., McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nguyen,T., Nicol,R., Norbu,C., O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P., Roman,J., Schauer,S., Schupback,R., Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M., Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.	
TITLE	Direct Submission	
JOURNAL	Submitted (03-JUN-2004) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA	
REFERENCE	8 (bases 1 to 214765)	
AUTHORS	Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N., Anderson,M., Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B., Camarata,J., Chang,J., Choepel,Y., Collymore,A., Cooke,A., Cooke,P., Corum,B., Dearellano,K., Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S., Graham,L., Grand-Pierre,N., Hafez,N., Hagopian,D., Hagos,B., Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K., Liu,G., Liu,X., Lui,A., Mabbitt,R., MacLean,C., Macdonald,P., Major,J., Manning,J., Matthews,C., McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nguyen,T., Nicol,R., Norbu,C., O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P., Roman,J., Schauer,S., Schupback,R., Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M., Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Z	

Fri Sep 9 11:21:43 2005

	AUTHORS	TITLE	JOURNAL	REFERENCE	AUTHORS
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repeat_region	13343. .13367	/rpt_family=" (TGGGG) n"			
repeat_region	14362. .14384				

Birren, B., Nusbaum, C. and Lander, E.
Mus musculus chromosome 7, clone RP23-152B12
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Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N.,
Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L.,
Boukhigal, B., Brown, A., Camarata, J., Campopiano, A., Chang, J.,
Chazaro, B., Choepel, Y., Colangelo, M., Collins, S., Collymore, A.,
Cook, A., Cooke, P., DeArelano, K., Dewar, K., Diaz, J. S., Dodge, S.,
Faro, S., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Gardyna, S.,
Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N.,
Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,
Kamat, A., Karatas, A., Kells, C., Larocque, K., Lamazares, R.,
Landers, T., Lehoczy, J., Levine, R., Lindblad-Toh, K., Liu, G.,
MacLean, C., Macdonald, P., Major, J., Marquis, N., Matthews, C.,
McCarthy, M., McEwan, P., McKernan, K., Meldrum, J., Meneus, L.,
Minhota, T., Mienga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R.,
Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neil, D.,
Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V.,
Raymond, C., Retta, R., Riback, M., Riley, R., Rise, C., Rogov, P.,
Roman, J., Roetti, M., Roy, A., Santos, R., Schauer, S., Schunback, R.,
Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
Strauss, N., Subramanian, A., Talamas, J., Testaye, S., Theodore, J.,
Topham, K., Travers, M., Travis, N., Trigglio, J., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G.,
Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE
JOURNAL
REFERENCE
AUTHORS

Direct Submission
Submitted (03-MAY-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 222540)
Birren, B., Nusbaum, C., Lander, E., Abouelleil, A., Allen, N., Anderson, M., Anderson, S., Arachchi, H.M., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhalter, B., Camarata, J., Chang, J., Choepel, Y., Collymore, A., Cook, A., Cooke, P., Corum, B., Dearallano, K., Diaz, J.S., Dodge, S., Dooley, K., Dorris, L., Erickson, J., Faro, S., Ferreira, P., FitzGerald, M., Gage, D., Hagalan, J., Gardyna, S., Graham, L., Grand-Pierre, N., Hafez, N., Hagopian, D., Hedges, B., Hall, J., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Katat, A., Karacas, A., Kellis, C., Landers, T., Levine, R., Lindblad-Toh, K., Liu, G., Liu, X., Lui, A., Mabbitt, R., Maclean, C., Macdonald, P., Major, J., Manning, J., Matthews, C., McCarthy, M., Meldrim, J., Menelus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nguyen, T., Nicol, R., Norbu, C., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Rachupka, A., Ramasamy, U., Raymond, C., Retta, R., Rice, C., Rogov, P., Roman, J., Schauer, S., Schupback, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Stubbs, M., Talamas, J., Tesfaye, S., Theodore, J., Toham, K., Travers, M., Vassiliev, H., Venkataraman, V.S., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.
Direct Submission
Submitted (16-SEP-2004) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Sep 16, 2004 this sequence version replaced gi:50284650.

TITLE	JOURNAL	COMMENT
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Zimmer, R. and Zody, W.
Direct Submission
Submitted (16-SEP-2004) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Sep 16, 2004 this sequence version replaced gi:50284650.
All repeats were identified using RepeatMasker:
Smit, A.P.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>
----- Genome Center
Center: Whitehead Institute/MIT Center for Genome Research
Center code: WIBR
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence_submissions@broad.mit.edu
----- Project Information
Center project name: L15655
Center Clone name: 152_B_12

* NOTE: This is a 'working draft' sequence. It currently
* consists of 7 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence

Query Match	84.8%	Score 17.8;	DB 10;	Length 214765;
Best Local Similarity	90.5%;	Pred. No. 3.9e+02;		
Matches 19;	Conservative 0;	Mismatches 2;	Indels 0;	Gaps 0;
QY	1	CAGTGACATGCAGGCTTAGCT 21		
Db	98096	CAGTGACTTCGAGGCTTAGCT 98116		
RESULT 20				
AC120123/c				
LOCUS	AC120123	222540 bp	DNA	linear
DEFINITION	Mus musculus chromosome 7 clone RP23-152B12 map 7,		***	SEQUENCING
	IN PROGRESS ***, 7 unordered pieces.			
ACCESSION	AC120123			
VERSION	AC120123.10	GI:52138883		
KEYWORDS	HTG; HTGS PHASE1; HTGS FULLTOP; HTGS_ACTIVEFIN.			
SOURCE	Mus musculus (house mouse)			
ORGANISM	Mus musculus			
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.			
REFERENCE	1	(bases 1 to 222540)		

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* as soon as it is available and the accession number will
* be preserved.
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* 47731: contig of 47731 bp in length
* 47732: gap of unknown length
* 47831: contig of 47831 bp in length
* 47832: gap of unknown length
* 61723: contig of 61723 bp in length
* 61724: gap of unknown length
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* 153132: gap of unknown length
* 153231: contig of 153231 bp in length
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* 170554: gap of unknown length
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* 190928: contig of 190928 bp in length
* 190929: gap of unknown length
* 216719: contig of 216719 bp in length
* 216720: gap of unknown length
* 222540: contig of 222540 bp in length.
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* Location/Qualifiers
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*     /mol_type="genomic DNA"
*     /db_xref="taxon:10090"
*     /chromosomes="7"
*     /map="7"
*     /clone="RP23-152B12"
*     /clone_lib="RPCI-23 Female Mouse BAC"
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* ORIGIN
*
* Query Match      84.8%; Score 17.8; DB 2; Length 222540;
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* Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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* QY 1 CAGTGACATGCAGGTCTAGCT 21
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* Db 177745 CAGTGATATGCAGGTCTAGCT 177725
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* RESULT 21
* AC115307
* LOCUS
* DEFINITION
* Rattus norvegicus clone CH230-11F18, WORKING DRAFT SEQUENCE.
* AC115307
* ACCESSION
* HTG: HTGS PHAS2; HTGS DRAFT: HTGS_FULLTOP.
* VERSION
* HTG: HTGS PHAS2; HTGS DRAFT: HTGS_FULLTOP.
* KEYWORDS
* Rattus norvegicus (Norway rat)
* SOURCE
* Rattus norvegicus
* Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
* Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
* Rattus.
*
* 1 (bases 1 to 260600)
* Muzny,D.Marie., Metzker,M.Lee., Abramson,S., Adams,C., Alder,J.,
* Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
* Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,B., Baden,H.,
* Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
* Biswal,K., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
* Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,
* Cardenas,V., Carter,K., Cavazos,I., Caesar,H., Center,A.,
* Chacko,J., Chavez,D., Chen,G., Chen,K., Chen,Y., Chen,Z., Chu,J.,
* Cleveland,C., Cockrell,R., Cox,C., Coyle,M., Cree,A., D'Souza,L.,
* Davila,M., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,
* Delgado,O., Denson,S., Deramo,C., Ding,Y., Dinh,H., Divya,K.,
* Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,
* Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G.,
* Fernandez,S., Finley,M., Flegg,N., Forbes,L., Foster,M., Foster,P.,
* Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,
* Gebregeorgis,E., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,W.,
* Gunaratne,P., Haaland,W., Hamil,C., Hamilton,N., Hamilton,K.,
* Harvey,Y., Havlak,P., Haves,A., Henderson,N., Hernandez,J.,
* Hernandez,R., Hines,S., Hladun,S.L., Hodgson,A., Hoques,M.,
* Hollins,B., Howells,S., Hulyk,S., Hume,J., Idlebird,D., Jackson,A.,
* Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,
* Karpathy,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
* Kowicz,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
* Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,
* Lorensueta,L., Loulesed,H., Lozado,R.J., Lu,X., Ma,J.,
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Mareshwari,M., Mahindartne,M., Mahmoud,M., Malloy,K., Mangum,A.,
Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E.,
Mawhiney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,
Milosavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,
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Nwaokelimeh,O., Okwunnu,G., Olarnpunaagoon,A., Pal,S., Parks,K.,
Pasternak,S., Paul,H., Perez,A., Perez,L., Pfannkuch,C.,
Plopper,F., Poindexter,A., Popovic,D., Primus,E., Pu,L., L.,
Puazo,M., Quiroz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R.,
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,
Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J.,
Sanders,W., Savery,G., Scherer,S., Scott,G., Shatsman,S., Shen,H.,
Shetty,J., Shvartsbeyn,A., Sisson,I., Sitter,C.D., Smajs,D.,
Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J.,
Staimle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Taylor,C.,
Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Umani,K.,
Valas,R., Vera,V., Villasana,D., Waldron,L., Walker,B., Wang,J.,
Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,
Williams,G., Willson,R., Wleczek,R., Wooden,H., Worley,K.,
Wright,D., Wright,J., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,
Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von
Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs,R.A.
Direct Submission
Unpublished
2 (bases 1 to 260600)
Worley,K.C.
Direct Submission
Submitted (17-MAR-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 260600)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (09-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Nov 9, 2002 this sequence version replaced gi:22772493.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GBBE
Center clone name: CH230-11F18
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 229014 bases at least Q40
Consensus quality: 231787 bases at least Q30
Consensus quality: 232904 bases at least Q20
Estimated insert size: 237320; sum-of-contigs estimation
Quality coverage: 7x in Q20 bases; sum-of-contigs estimation
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* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 1 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
```


* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 1 contigs. Gaps between the contigs
 * are represented as runs of N. The order of the pieces
 * is believed to be correct as given, however the sizes
 * of the gaps between them are based on estimates that have
 * provided by the submitter.

* This sequence will be replaced
 * by the finished sequence as soon as it is available and
 * the accession number will be preserved.
 * 1 281447: contig of 281447 bp in length.

FEATURES

source
 1..281447
 /organism="Rattus norvegicus"
 /mol_type="genomic DNA"
 /db_xref="taxon:10116"
 /clone="CH230-233D15"
 misc_feature
 1..1083
 /note="wgs contig"
 misc_feature
 2987..4464
 /note="wgs contig"
 misc_feature
 192314..192414
 /note="clone boundary
 clone_end:Sp6
 site:5cORI
 end_sequence:B2091967"
 280368..281447
 /note="wgs end-extension
 clone_end:Sp6"

ORIGIN

Query Match 84.8%; Score 17.8; DB 2; Length 281447;

Best Local Similarity 90.5%; Pred. No. 3.8e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTTAGCT 21

Db 264359 CTGTGATATGCAGGCTTAGCT 264339

RESULT 23

BD185177/c 5446 bp DNA linear PAT 17-JUN-2003
 LOCUS
 DEFINITION Novel genes and proteins encoded by the genes.

ACCESSION BD185177.1 GI:31877377

VERSION JP 2002345493-A/20.

KEYWORDS Homo sapiens (human)

SOURCE

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS Ohara,O., Nagase,T. and Nakajima,D.

TITLE Novel genes and proteins encoded by the genes

JOURNAL Patent: JP 2002345493-A 20 03-DEC-2002;

COMMENT KAZUSA DNA RESEARCH INSTITUTE

OS Homo sapiens (human)

PN JP 2002345493-A/20

PD 03-DEC-2002

PF 26-FEB-2002 JP 2002049046

PI OSAMU OHARA, TAKAHIRO NAGASE, DAISUKE NAKAJIMA

PC C12N15/09,C07K14/47,C07K14/54,C12N15/00

CC Novel genes and proteins encoded by the genes FH Key

Location/Qualifiers

FT CDS (2564)..(4138).

Location/Qualifiers

1..5446

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

ORIGIN

Query Match 82.9%; Score 17.4; DB 6; Length 5446;

Best Local Similarity 94.7%; Pred. No. 9.5e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTTAG 19

Db 833 CAGTGACAGGCAGGCTTAG 815

RESULT 24

AC121561/c

LOCUS

DEFINITION

AC121561

AC121561.1 GI:20986629

HTG: HTGS_PHASE0.

SOURCE Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS Birren,B., Linton,L., Nusbaum,C. and Lander,E.

JOURNAL Unpublished

REFERENCE

2 (bases 1 to 46070)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,

Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavsky,L.,

Boukhgalter,B., Brown,A., Camarata,J., Campopiano,A., Chang,J.,

Chazaro,B., Choepel,Y., Collangelo,M., Collins,S., Collymore,A.,

Cook,A., Cooke,P., DeAtrellano,K., Dewar,K., Diaz,J.S., Dodge,S.,

Faro,S., Ferreira,P., Fitzgerald,M., FitzHugh,W., Gage,D.,

Galagan,J., Gardyna,S., Ginde,S., Gord,S., Goyette,M., Graham,L.,

Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I.,

Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., LaRoque,K.,

Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Lindblad-Toh,K.,

Liu,G., MacLean,C., MacDonald,P., Major,J., Marquis,N.,

Mathews,C., McCarthy,M., McEwan,P., McKernan,K., Meldrum,J.,

Meneus,L., Mihova,T., Mienga,V., Murphy,T., Naylor,J., Nguyen,C.,

Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P.,

O'Neil,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,

Pollara,V., Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C.,

Rogov,P., Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S.,

Schupbach,R., Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N.,

Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S.,

Theodore,J., Topham,K., Travers,M., Travis,N., Trigilio,J.,

Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,

Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

Direct Submission

Submitted (20-MAY-2002) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

All repeats were identified using RepeatMasker:

Smit, A.F.A. & Green, P. (1996-1997)

<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: <http://www-seq.wi.mit.edu>

Contact: sequence_submission@genome.wi.mit.edu

Center project name: L26773

Center clone name: 2527_E_13

* NOTE: This record contains 58 individual
 * sequencing reads that have not been assembled into
 * contigs. Runs of N are used to separate the reads
 * and the order in which they appear is completely
 * arbitrary. Low-pass sequence sampling is useful for
 * identifying clones that may be gene-rich and allows
 * overlap relationships among clones to be deduced.
 * However, it should not be assumed that this clone
 * will be sequenced to completion. In the event that
 * the record is updated, the accession number will
 * be preserved.

29403: gap of 100 bp
30080: contig of 677 bp in length
30180: gap of 100 bp
30844: contig of 664 bp in length
30944: gap of 100 bp
31640: contig of 696 bp in length
31740: gap of 100 bp
32459: contig of 719 bp in length
32559: gap of 100 bp
33262: contig of 703 bp in length
33362: gap of 100 bp
34090: contig of 728 bp in length
34190: gap of 100 bp
34892: contig of 702 bp in length
34992: gap of 100 bp
35692: contig of 700 bp in length
35792: gap of 100 bp
36480: contig of 688 bp in length
36580: gap of 100 bp
37293: contig of 713 bp in length
37393: gap of 100 bp
38091: contig of 698 bp in length
38191: gap of 100 bp
38897: contig of 706 bp in length
38997: gap of 100 bp
39720: contig of 723 bp in length
39820: gap of 100 bp
40526: contig of 706 bp in length
40626: gap of 100 bp
41302: contig of 676 bp in length
41402: gap of 100 bp
42077: contig of 675 bp in length
42177: gap of 100 bp
42841: contig of 664 bp in length
42941: gap of 100 bp
43738: gap of 100 bp
44456: contig of 718 bp in length
44556: gap of 100 bp
45255: contig of 699 bp in length
45355: gap of 100 bp
46070: contig of 715 bp in length.

29304
29404
30081
30181
30845
30945
31641
31741
32460
32560
33263
33363
34091
34191
34893
34993
35693
35793
36481
36581
37294
37394
38092
38192
38898
38998
39721
39821
40527
40627
41303
41403
42078
42178
42842
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43739
44457
44557
45256
45356
46070

Location/Qualifiers
1. .46070
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="17"
/map="17"
/clone="CTD-2527E13"
/clone_lib="CITD2 Human BAC"

Query Match 82.9%; Score 17.4; DB 2; Length 46070;
Best Local Similarity 94.7%; Pred. No. 7.4e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTAG 19
|||||
Db 31187 CAGTGACATGCAGGCTAG 31169

RESULT 25
AL133227
LOCUS
DEFINITION
Human DNA sequence from clone RP11-39402 on chromosome 20. Contains the gene for CGI-15 protein, a gene for a novel protein similar to KIAA0281 and Drosophila CG5336, ESTs, STSs, GSSs and a CpG island, complete sequence.
ACCESSION
AL133227
VERSION
AL133227.15 GI:9187135
KEYWORDS
HTG; CpG island; KIAA0281.
SOURCE
Homo sapiens (human)

684: contig of 684 bp in length
784: gap of 100 bp
1401: contig of 617 bp in length
1501: gap of 100 bp
2203: contig of 702 bp in length
2303: gap of 100 bp
2984: contig of 681 bp in length
3084: gap of 100 bp
3754: contig of 670 bp in length
3854: gap of 100 bp
4545: contig of 691 bp in length
4645: gap of 100 bp
5346: contig of 701 bp in length
5446: gap of 100 bp
6142: contig of 696 bp in length
6242: gap of 100 bp
6949: contig of 707 bp in length
7049: gap of 100 bp
7730: contig of 681 bp in length
7830: gap of 100 bp
8515: contig of 685 bp in length
8615: gap of 100 bp
9267: contig of 652 bp in length
9367: gap of 100 bp
10068: contig of 701 bp in length
10168: gap of 100 bp
10859: contig of 691 bp in length
10959: gap of 100 bp
11667: contig of 708 bp in length
11767: gap of 100 bp
12472: contig of 705 bp in length
12572: gap of 100 bp
13277: contig of 705 bp in length
13377: gap of 100 bp
14069: contig of 692 bp in length
14169: gap of 100 bp
14870: contig of 701 bp in length
14970: gap of 100 bp
15681: contig of 711 bp in length
15781: gap of 100 bp
16482: contig of 701 bp in length
16582: gap of 100 bp
17271: contig of 689 bp in length
17371: gap of 100 bp
18068: contig of 697 bp in length
18168: gap of 100 bp
18863: contig of 695 bp in length
18963: gap of 100 bp
19650: contig of 687 bp in length
19750: gap of 100 bp
20456: contig of 706 bp in length
20556: gap of 100 bp
21267: contig of 711 bp in length
21367: gap of 100 bp
22078: contig of 711 bp in length
22178: gap of 100 bp
22875: contig of 697 bp in length
22975: gap of 100 bp
23661: contig of 686 bp in length
23761: gap of 100 bp
24488: contig of 727 bp in length
24588: gap of 100 bp
25276: contig of 688 bp in length
25376: gap of 100 bp
26082: contig of 706 bp in length
26182: gap of 100 bp
26886: contig of 704 bp in length
26986: gap of 100 bp
27691: contig of 705 bp in length
27791: gap of 100 bp
28494: contig of 703 bp in length
28594: gap of 100 bp
29303: contig of 709 bp in length

1
685
785
1402
1502
2204
2304
2985
3085
3755
3855
4546
5347
5447
6143
6243
6950
7050
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9368
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11668
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12573
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14871
14971
15682
15782
16483
16583
17272
17372
18069
18169
18864
18964
19651
19751
20457
20557
21268
21368
22079
22179
22876
22976
23662
23762
24489
24589
25277
25377
26083
26183
26887
26987
27692
27792
28495
28595

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 85566)

REFERENCE
AUTHORSTITLE
JOURNAL

Direct Submission
Submitted (02-APR-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
requests: clonerequest@sanger.ac.uk

COMMENT

On Jul 14, 2000 this sequence version replaced gi:8977995.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome 20, constructed by the Sanger Centre Chromosome 20
Mapping Group. Further information can be found at
<http://www.sanger.ac.uk/HGP/Chr20>
IMPORTANT: This sequence is not the entire insert of clone
RP11-39402 it may be shorter because we sequence overlapping
sections only once, except for a 100 base overlap.
The true left end of clone RP11-39402 is at 1 in this sequence. The
true left end of clone RP5-981123 is at 85467 in this sequence. The
true right end of clone RP5-984123 is at 78450 in this sequence.
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest. RP11-39402 is from
the library RPCI-11.2 constructed by the group of Pieter de Jong.
For further details see
<http://www.chori.org/bacpac/home.htm>
VECTOR: pBACe3.6.

FEATURES

Source

Location/Qualifiers

1..85566
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="20"
/clone="RP11-39402"
/clone_lib="RPCI-11.2"
33..90
/notes="MIR repeat: matches 107. .161 of consensus"
139..438
/notes="AluX repeat: matches 1. .293 of consensus"
1404..1535
/notes="MIR repeat: matches 54. .182 of consensus"
3358..3536
/notes="L2 repeat: matches 2318. .2500 of consensus"
3438..3969
/notes="match: GSS: Em:AQ373404"
3456..3976
/notes="match: GSS: Em:AQ885421"
3620..3905
/notes="MIR repeat: matches 16. .262 of consensus"
3653..4183
/notes="match: GSS: Em:AQ374823"
5271..5327
/notes="MIR repeat: matches 61. .121 of consensus"
6122..6156
/notes="MIR repeat: matches 110. .144 of consensus"
6577..6685
/notes="L1M4 repeat: matches 5209. .5334 of consensus"
7049..7178

repeat_region

repeat_region

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repeat_region

misc_feature

misc_feature

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repeat_region

/note="L2 repeat: matches 2562. .2695 of consensus"
7275..7472

/note="MIR repeat: matches 44. .254 of consensus"
7545..7917

/note="ITR16C repeat: matches 11. .381 of consensus"
7733..8207

/note="match: GSS: Em:AQ524007"
8425..8510

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8622..8835

/note="MIR repeat: matches 35. .262 of consensus"
8945..9114

/note="MIR repeat: matches 21. .210 of consensus"
9536..9599

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9666..9851

/note="MIR repeat: matches 80. .262 of consensus"
10546..10783

/note="match: GSS: Em:AQ543754"
11352..11438

/note="MIR repeat: matches 85. .168 of consensus"
12303..12620

/note="match: GSS: Em:AQ195196
match: STS: Em:G52843"
12324..12752

/note="match: GSS: Em:AQ631329"
12855..13015

/note="MIR repeat: matches 65. .220 of consensus"
13723..14037

/note="AluDb repeat: matches 16. .312 of consensus"
complement(13988..14650)

/note="match: GSS: Em:AQ899717"
complement(14441..14659)

/note="match: GSS: Em:AQ62441"
15533..15742

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16607..16778

/note="MER91A repeat: matches 2. .171 of consensus"
17496..17568

/note="L2 repeat: matches 2624. .2696 of consensus"
17746..18049

/note="AluX repeat: matches 1. .304 of consensus"
18067..18413

/note="L1P413 repeat: matches 5807. .6156 of consensus"
18593..18825

/note="MIR repeat: matches 20. .248 of consensus"
18982..19111

/note="MIR repeat: matches 136. .250 of consensus"
19112..19406

/note="AluDb repeat: matches 1. .296 of consensus"
19407..19518

/note="MIR repeat: matches 20. .136 of consensus"
20100..20283

/note="L2 repeat: matches 2550. .2750 of consensus"
20312..20763

/note="Charlie4 repeat: matches 17. .485 of consensus"
20787..21340

/note="L2 repeat: matches 1936. .2534 of consensus"
21447..21736

/note="MLTJ repeat: matches 1. .320 of consensus"
21990..22289

/note="AluY repeat: matches 1. .301 of consensus"
22377..22462

/note="L2 repeat: matches 2621. .2710 of consensus"
22463..22749

/note="AluX repeat: matches 1. .292 of consensus"
24255..24319

/note="MIR repeat: matches 109. .175 of consensus"
24663..24768

/note="MIR repeat: matches 80. .183 of consensus"
24832..24928

/note="MIR repeat: matches 92. .190 of consensus"
26048..26173

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26722. .26765
repeat_region /note="11 copies 4 mer catc 93% conserved"
26766. .26809
repeat_region /note="11 copies 4 mer atcc 81% conserved"
26813. .26844
repeat_region /note="8 copies 4 mer atcc 93% conserved"
26839. .26918
repeat_region /note="4 copies 20 mer 77% conserved"
27222. .27559
repeat_region /note="LTR16A repeat: matches 95. .440 of consensus"
27583. .27748
repeat_region /note="FRAM repeat: matches 1. .166 of consensus"
27786. .27897
repeat_region /note="L2 repeat: matches 2637. .2750 of consensus"
28308. .28431
repeat_region /note="Alusq repeat: matches 1. .132 of consensus"
28448. .28524
repeat_region /note="LTR16A repeat: matches 193. .270 of consensus"
28525. .28714
repeat_region /note="Alusq repeat: matches 114. .301 of consensus"
28837. .29017
repeat_region /note="MIR repeat: matches 65. .250 of consensus"
29053. .29187
repeat_region /note="MIR repeat: matches 84. .226 of consensus"
30560. .31063
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30596. .30931
misc_feature /note="match: GSS: Em:AQ901120"
30602. .30764
repeat_region /note="MER58A repeat: matches 46. .221 of consensus"
complement(31347. .46204)
/genes="BA39402.1"
mRNA complement(join(31347. .32333,32570. .32699,33860. .34035,
36691. .36774,36882. .37050,37611. .37683,38383. .38446,
39432. .39582,40196. .40572,46180. .46204))
/genes="BA39402.1"
/product="BA39402.1 (CGI-15 protein)"
/note="match: cDNAs: Em:AF132949
match: ESTs: Em:AU079917 Em:AA667893 Em:AI007286
Em:AU067617 Em:W56183 Em:R69763 Em:H06603 Em:AW631237

```

Query Match 82.9%; Score 17.4; DB 9; Length 85566;
 Best Local Similarity 94.7%; Pred. No. 6.9e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATCGAGTCTAG 19
 Db 64014 CAGTGACAGCGAGTCTAG 64032

Search completed: September 6, 2005, 20:29:56
 Job time : 746.656 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 17:45:55 ; Search time 1500.84 Seconds
(without alignments)
532.600 Million cell updates/sec

Title: US-10-729-421-40

Perfect score: 21
Sequence: 1 cagtgcacatgcaggttagct 21

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 100 summaries

Database :

EST: *
1: gb_est1: *
2: gb_est2: *
3: gb_hic: *
4: gb_est3: *
5: gb_est4: *
6: gb_est5: *
7: gb_est6: *
8: gb_gsa1: *
9: gb_gsa2: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	19	90.5	1041	9	CNS004CAJ
2	18.4	87.6	1157	6	CD500170
3	17.8	84.8	211	8	AZ995872
4	17.8	84.8	611	8	AZ218630
5	17.8	84.8	878	9	CR794665
6	17.8	84.8	4261	3	AK083880
7	17.4	82.9	380	9	CL289600
8	17.4	82.9	718	9	CG176005
9	17.4	82.9	722	2	BF607360
10	17.4	82.9	772	8	CC340825
11	17.4	82.9	772	9	CG211386
12	17.4	82.9	805	9	CG362883
13	17.4	82.9	816	4	B1548151
14	17.4	82.9	817	8	CC340835
15	17.4	82.9	923	9	CG211861
16	17	81.0	382	6	BY609340
17	17	81.0	408	6	BY649695
18	17	81.0	622	6	CA124343
19	17	81.0	633	8	AZ069535
20	17	81.0	642	2	BB650662
21	17	81.0	655	2	BB233162
22	17	81.0	768	9	AG457137
23	17	81.0	781	9	AG557182
24	17	81.0	949	7	CF411086

AK044974	Mus muscu	AK044974	2445	81.0	17	3	AK044974
AK013040	Mus muscu	AK013040	2466	81.0	17	3	AK013040
AK017012	Mus muscu	AK017012	2559	81.0	17	3	AK017012
AK082079	Mus muscu	AK082079	3256	81.0	17	3	AK082079
AK081942	Mus muscu	AK081942	3417	81.0	17	3	AK081942
AW846933	RC3-C7019	AW846933	306	80.0	16.8	2	AW846933
AU249500	AU249500	AU249500	330	80.0	31	1	AU249500
AA050319	mjl1405.f	AA050319	335	80.0	32	1	AA050319
BY077663	BY077663	BY077663	387	80.0	33	5	BY077663
AA004028	mg80G02.f	AA004028	400	80.0	34	1	AA004028
AA117056	ml29C06.f	AA117056	400	80.0	35	1	AA117056
AA003782	mg2e10.f	AA003782	407	80.0	36	1	AA003782
BE656207	UI-M-BH0-	BE656207	407	80.0	37	2	BE656207
AA052553	mc66D04.f	AA052553	414	80.0	38	1	AA052553
W79973	me90e09.f	W79973	419	80.0	39	7	W79973
AA008849	mg98F08.f	AA008849	426	80.0	40	1	AA008849
BY273688	BY273688	BY273688	435	80.0	41	5	BY273688
BY240303	BY240303	BY240303	436	80.0	42	5	BY240303
BY051479	BY051479	BY051479	442	80.0	43	5	BY051479
CB788954	AMGNNUC:M	CB788954	443	80.0	44	6	CB788954
BX529253	BX529253	BX529253	461	80.0	45	5	BX529253
W89580	mf73f08.f	W89580	466	80.0	46	7	W89580
AJ647590	AJ647590	AJ647590	467	80.0	47	1	AJ647590
BB863212	BB863212	BB863212	467	80.0	48	2	BB863212
AJ647892	AJ647892	AJ647892	469	80.0	49	1	AJ647892
CA533118	C0345E08-	CA533118	508	80.0	50	6	CA533118
W89380	mf73g08.f	W89380	510	80.0	51	7	W89380
BF556581	UI-R-E1-f	BF556581	512	80.0	52	2	BF556581
BE374887	601226811	BE374887	515	80.0	53	2	BE374887
BZ132704	CH230-481	BZ132704	523	80.0	54	8	BZ132704
AA712019	UI-M-HA0-	AA712019	542	80.0	55	1	AA712019
CF732719	UI-M-HA0-	CF732719	544	80.0	56	7	CF732719
BH347218	CH230-42D	BH347218	546	80.0	57	8	BH347218
BG276278	uv02610.y	BG276278	549	80.0	58	4	BG276278
AA682125	vu13003.f	AA682125	552	80.0	59	1	AA682125
AA530593	VJ49G09.f	AA530593	561	80.0	60	1	AA530593
BF452552	mao1d05.f	BF452552	568	80.0	61	2	BF452552
AV597290	AV597290	AV597290	570	80.0	62	1	AV597290
AV595391	AV595391	AV595391	576	80.0	63	1	AV595391
AL792667	AL792667	AL792667	582	80.0	64	1	AL792667
BQ840540	mah69f08	BQ840540	584	80.0	65	5	BQ840540
CG672097	RRM266 Ba	CG672097	588	80.0	66	9	CG672097
BG100744	uy14c01.y	BG100744	596	80.0	67	4	BG100744
BI985696	3144-63 M	BI985696	600	80.0	68	4	BI985696
AW412020	uo55h02.y	AW412020	617	80.0	69	2	AW412020
CR421241	CR421241	CR421241	619	80.0	70	1	CR421241
CK621976	ml31a12.y	CK621976	624	80.0	71	7	CK621976
BJ774417	BJ774417	BJ774417	626	80.0	72	4	BJ774417
CN119882	ECOCAA002	CN119882	627	80.0	73	7	CN119882
BG099869	uy13c02.y	BG099869	637	80.0	74	4	BG099869
CD766336	AGENCOURT	CD766336	641	80.0	75	6	CD766336
AL863627	AL863627	AL863627	642	80.0	76	1	AL863627
AL878478	AL878478	AL878478	642	80.0	77	1	AL878478
CF732690	UI-M-HA0-	CF732690	649	80.0	78	7	CF732690
BX315194	BX315194	BX315194	651	80.0	79	5	BX315194
CD772714	AGENCOURT	CD772714	655	80.0	80	1	CD772714
AZ428022	IM0210N17	AZ428022	660	80.0	81	8	AZ428022
BG381620	UI-R-CT0-	BG381620	679	80.0	82	4	BG381620
AG151855	pan trogl	AG151855	685	80.0	83	9	AG151855
BQ746626	UI-M-ER0-	BQ746626	688	80.0	84	5	BQ746626
BK654484	AGENCOURT	BK654484	701	80.0	85	7	BK654484
AZ247538	RPCI-23-9	AZ247538	720	80.0	86	8	AZ247538
AZ247542	RPCI-23-9	AZ247542	720	80.0	87	8	AZ247542
CF148724	AGENCOURT	CF148724	734	80.0	88	7	CF148724
COB06312	AGENCOURT	COB06312	734	80.0	89	7	COB06312
CB598900	AGENCOURT	CB598900	738	80.0	90	6	CB598900
AG527299	Mus muscu	AG527299	749	80.0	91	6	AG527299
CO362201	AGENCOURT	CO362201	753	80.0	92	7	CO362201
CR441613	CR441613	CR441613	753	80.0	93	7	CR441613
CF149901	AGENCOURT	CF149901	765	80.0	94	7	CF149901
CC553492	CH240_459	CC553492	768	80.0	95	9	CC553492
AY413996	Mus muscu	AY413996	822	80.0	96	1	AY413996
BU412600	602954786	BU412600	829	80.0	97	5	BU412600

98 16.8 80.0 835 4 BI684946 BI684946 603307795
 C 99 16.8 80.0 844 9 CR017849 Forward s
 100 16.8 80.0 847 7 CK793898 AGENCOURT

ALIGNMENTS

RESULT 1
 LOCUS CNS04CAJ
 DEFINITION Tetraodon nigroviridis genome survey sequence T7 end of clone
 099K23 of library G from Tetraodon nigroviridis, genomic survey
 sequence.

ACCESSION AL284212
 VERSION AL284212.1 GI:8022590
 KEYWORDS GSS; genome survey sequence.
 SOURCE Tetraodon nigroviridis
 ORGANISM Tetraodon nigroviridis

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
 Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
 Tetraodontidae; Tetraodontidae; Tetraodon.

REFERENCE 1
 AUTHORS Roest Crolius,H., Jaillon,O., Dasilva,C., Bouneau,L., Fisher,C.,
 Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
 Saurin,W. and Weissenbach,J.
 TITLE Estimate of human gene number provided by genome-wide analysis
 using Tetraodon nigroviridis DNA sequence
 JOURNAL Nat. Genet. 25 (2), 235-238 (2000)
 MEDLINE 20296633
 PUBMED 10835645

REFERENCE 2
 AUTHORS Roest Crolius,H., Jaillon,O., Dasilva,C., Ozouf-Costaz,C.,
 Fizames,C., Fischer,C., Bouneau,L., Billault,A., Quetier,F.,
 Saurin,W., Bernot,A. and Weissenbach,J.
 TITLE Characterization and repeat analysis of the compact genome of the
 freshwater pufferfish Tetraodon nigroviridis
 JOURNAL Genome Res. 10 (7), 939-949 (2000)
 MEDLINE 20359837
 PUBMED 10899143

REFERENCE 3 (bases 1 to 1041)
 AUTHORS Genoscope.
 TITLE Direct Submission
 JOURNAL Submitted (12-APR-2000) Genoscope - Centre National de Sequencage :
 BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
 - Web : www.genoscope.cns.fr)

COMMENT This sequence is a single read and was generated as part of a large
 scale clone-end sequencing project of the Tetraodon nigroviridis
 genome. For more information, please take a look at
 http://www.genoscope.cns.fr/Tetraodon.

FEATURES

source

1..1041
 /location="Tetraodon nigroviridis"
 /mol_type="genomic DNA"
 /db_xref="taxon:99883"
 /clone="099K23"
 /clone_lib="G"
 /notes="Genoscope sequence ID : C0BG099AF12LP1-end : T7"

ORIGIN

Query Match 90.5%; Score 19; DB 9; Length 1041;
 Best Local Similarity 90.5%; Pred. No. 2e+02;
 Matches 19; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCTAGCT 21
 :|||||

Db 942 SAGTGACATGCAGGTCTACCT 962
 :|||||

RESULT 2
 CD500170
 LOCUS

DEFINITION
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM

CDA43-E06_x1d-t_SHGC-CDA Gasterosteus aculeatus cDNA clone
 CDA43-E06 5', mRNA sequence.
 CD500170
 CD500170.1 GI:31427201
 EST.

Gasterosteus aculeatus (three spined stickleback)

Gasterosteus aculeatus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
 Acanthomorpha; Acanthopterygii; Percomorpha; Gasterosteiformes;
 Gasterosteidae; Gasterosteus.

REFERENCE 1 (bases 1 to 1157)
 AUTHORS Kingsley,D.M., Peichel,C., Balabhadra,S., Grimwood,J., Dickson,M.,
 Schmutz,J. and Myers,R.M.
 TITLE Expressed sequence tags from Gasterosteus aculeatus
 JOURNAL Unpublished (2003)
 COMMENT Contact: Kingsley, DM
 HHMI and Department of Developmental Biology
 Stanford University School of Medicine
 Beckman Center B300, 279 Campus Drive, Stanford, CA 94305-5329, USA
 Tel: 650 725 5954
 Fax: 650 725 7739
 Email: kingsley@cmgm.stanford.edu
 Plate: 43

High quality sequence stop: 785.

FEATURES
 source

1..1157
 /location="Gasterosteus aculeatus"
 /mol_type="mRNA"
 /strain="Salinas river, CA"
 /db_xref="taxon:69293"
 /clone="CDA43-E06"
 /sex="mixed male and female"
 /tissue_type="heads and internal organs combined"
 /dev_stage="adult"
 /clone_lib="SHGC-CDA"
 /note="Vector: lambda ZAP Express/pBK-CMV; Site 1: EcoRI
 (5' adaptor); Site 2: XhoI (3' linker primer); The mixed
 organ cDNA library was generated using the ZAP-CDNA method
 by Stratagene. First strand cDNA synthesis was primed with
 a 50 bp linker primer containing an oligo dt sequence
 preceded by a synthetic XhoI site. 5 prime adaptors were
 used containing an EcoRI cohesive end. The finished cDNAs
 were inserted in to the ZAP express vector
 unidirectionally in the sense orientation with respect to
 the lacZ promoter of pBK-CMV. An amplified library was
 prepared from approximately 3 million primary clones in
 the lambda ZAP Express vector. In vivo excision was then
 used to generate individual pBK-CMV phagemid clones for
 EST sequencing."

ORIGIN

Query Match 87.6%; Score 18.4; DB 6; Length 1157;
 Best Local Similarity 95.0%; Pred. No. 4e+02;
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 2 AGTGACATGCAGGTCTAGCT 21
 :|||||
 Db 132 AGTGACATGCAGGTCTACCT 151

RESULT 3
 AZ995872
 LOCUS

DEFINITION
 AZ995872 211 bp DNA linear GSS 27-APR-2001
 2M0281B23R Mouse 10kb plasmid UUGC2M library Mus musculus genomic
 clone UUGC2M0281B23 R, genomic survey sequence.

ACCESSION AZ995872
 VERSION AZ995872
 KEYWORDS GSS.
 SOURCE Mus musculus (house mouse)

ORGANISM

Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 211)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Iglam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0281 row: B column: 23
 Seq primer: CACACGGAACAGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 211.

FEATURES

source

Location/Qualifiers

1..211
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC2M0281B23"
 /sex="Female"
 /lab_host="E. coli strain XL10-Gold, Tl-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUGC2M library"
 /notes="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (female) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (GI|4732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

ORIGIN

Query Match 84.8%; Score 17.8; DB 8; Length 211;
 Best Local Similarity 90.3%; Pred. No. 6.1e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTTAGCT 21
 ||||| ||||| ||||| ||||| |||||

Db 136 CAGAGCCATGCAGGCTTAGCT 156

RESULT 4

AZ218630/c

LOCUS AZ218630 611 bp DNA linear GSS 09-JUN-2000
 DEFINITION Sheared DNA-82B8.TF Sheared DNA Trypanosoma brucei genomic clone

ACCESSION AZ218630
 VERSION AZ218630.1 GI:8436430

KEYWORDS GSS.

SOURCE

Trypanosoma brucei

Trypanosoma brucei

Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;

Trypanosoma.

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

1 (bases 1 to 611)
 El-Sayed,N., Zhao,S., Zhao,H., Gill,S., Suh,E., Malek,J., Fujii,C., Gerard,C., Leech,V., de Jong,P., Ullu,E., Melville,S., Donelson,J., Frazer,C. and Adams,M.
 Determination of clone end sequences from Trypanosoma brucei GUTat 10.1 sheared DNA library
 Unpublished (1999)
 Other GSSs: Sheared DNA-82B8.TR
 Contact: Najib M. El-Sayed
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: nelsayed@tigr.org
 Clones are derived from the Trypanosoma brucei GUTat 10.1 sheared DNA library constructed at TIGR. Clones will be available for distribution through Research Genetics, Alabama, USA. Sheared DNA end sequences search page: <http://www.tigr.org/tdb/mdb/tbdb/>.
 Seq primer: M13-Forward
 Class: shotgun.

FEATURES

source

Location/Qualifiers

1..611
 /organism="Trypanosoma brucei"
 /mol_type="genomic DNA"
 /strain="TREU927/4 GUTat 10.1"
 /db_xref="taxon:5691"
 /clone="Sheared DNA-82B8"
 /clone_lib="Sheared DNA"
 /note="Vector: pUC18; Site1: SmaI; Constructed at The Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of Trypanosoma brucei (TREU927/4 GUTat 10.1) was mechanically sheared to give a tight size distribution (approx 2 kb). The v + i method used for the library construction is described in detail in Smith, H.O. and Venter, J.C. (Making small insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaudin and B. Barrell, Oxford University Press, 1999)."

ORIGIN

Query Match 84.8%; Score 17.8; DB 8; Length 611;
 Best Local Similarity 90.5%; Pred. No. 7.2e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTTAGCT 21
 ||||| ||||| ||||| ||||| |||||

Db 548 CAGTGGCATGCAGGCTTAGTT 528

RESULT 5

CR794665

LOCUS

DEFINITION GROAA12AC07RM1 INRA BAC Bos taurus genomic clone INRA_B225E12, DNA sequence, genomic survey sequence.

ACCESSION

CR794665

VERSION

CR794665.1

KEYWORDS

GSS.

SOURCE

Bos taurus (cow)

Bos taurus

Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;

Bovinae; Bos.

REFERENCE 1 (bases 1 to 878)

AUTHORS

Eggen,A., Schibler,L. and Roy,A.

TITLE

Bovine BAC End Sequences from the INRA bovine BAC library

JOURNAL

Unpublished

REFERENCE 2 (bases 1 to 878)

Genoscope.

Direct Submission

TITLE

Submitted (20-SEP-2004) Genoscope - Centre National de Sequençage :

JOURNAL

BP 191 91006 EVRY cedex - FRANCE (E-mail : segref@genoscope.cns.fr)

```

- Web : www.genoscope.cns.fr)
Contact: Andre Eggen
Department of Animal Genetics - LGbC
INRA
78350 Jouy-en-Josas, France
Tel: 33 1 34 65 24 24
Fax: 33 1 34 65 24 78
Email: eggen@jouy.inra.fr
Clones are derived from the INRA bovine BAC library
(http://locus.jouy.inra.fr/fpc/cattle_bac_map.htm). For BAC library
availability, please contact Andre Eggen (eggen@jouy.inra.fr). This
work was undertaken as part of the International Bovine BAC
Mapping Consortium (IBBMC) by INRA (Jouy-en-Josas) and Genoscope
(Evry) Plate: 225 row: B column: 12
Seq primer: M13 Reverse
Class: BAC ends.

FEATURES
Source
1..878
/organism="Bos taurus"
/mol_type="genomic DNA"
/strain="breed: Holstein"
/db_xref="taxon:9913"
/clone="INRab_225E12"
/sex="Male"
/cell_type="fibroblast"
/clone_lib="INRA bovine BAC"
/notes="Vector: pBeloBAC11; Site_1: HindIII; Holstein bull;
INRA Bovine BAC library (Male) produced by Andre
Eggen-Genoscope sequence ID : GR0AAA12AC07RM1"

ORIGIN
Query Match 84.8%; Score 17.8; DB 9; Length 878;
Best Local Similarity 90.5%; Pred. No. 7.7e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTTAGCT 21
|||||
Db 297 CAGTCACAGCAGGCTTAGCT 317

RESULT 6
AK083880/c
LOCUS
DEFINITION
Mus musculus 12 days embryo spinal ganglion cDNA, RIKEN full-length
enriched library, clone: D130043C18 product: unclassifiable, full
insert sequence.
ACCESSION
AK083880
VERSION
AK083880.1 GI:26101557
KEYWORDS
HTC; CAP trapper.
SOURCE
Mus musculus (house mouse)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1
Carninci, P. and Hayashizaki, Y.
2
High-efficiency full-length cDNA cloning
Meth. Enzymol. 303, 19-44 (1999)
99279253
10349636
PUBMED
REFERENCE
1
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new genes
Genome Res. 10 (10), 1617-1630 (2000)
20493374
11042159
PUBMED
REFERENCE
2
Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
Konno, H., Akiyama, J., Nishi, K., Kitsuunai, T., Tashiro, H., Itoh, M.,
Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A.,
Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K.,
Fujiwaki, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M.,
Yoneda, Y., Iehikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J.,
Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multicapillary sequencer
Genome Res. 10 (11), 1757-1771 (2000)
20530913
11076861
PUBMED
REFERENCE
4
The RIKEN Genome Exploration Research Group Phase II Team and the
FANTOM Consortium.
Functional annotation of a full-length mouse cDNA collection
Nature 409, 685-690 (2001)
JOURNAL
NATURE
AUTHORS
5
The FANTOM Consortium and the RIKEN Genome Exploration Research
Group Phase I & II Team.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)
JOURNAL
NATURE
REFERENCE
6
(bases 1 to 4261)
Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P.,
Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W.,
Hayashida, K., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T.,
Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kasukawa, T.,
Kato, H., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Kouda, M.,
Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, M.,
Nakamura, M., Nishi, K., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N.,
Okazaki, Y., Saito, R., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N.,
Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T.,
Sogabe, Y., Tagami, M., Tagawa, A., Takahashi, F., Takaku-Akahira, S.,
Takeda, Y., Tanaka, T., Tomaru, A., Toya, T., Yasunishi, A.,
Muramatsu, M. and Hayashizaki, Y.
Direct Submission
Submitted (16-APR-2002) Yoshihide Hayashizaki, The Institute of
Physical and Chemical Research (RIKEN), Laboratory for Genome
Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
RIKEN Yokohama Institute, 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
Kanagawa 230-0045, Japan (E-mail: genome-res@sc.riken.jp,
URL: http://genome.gsc.riken.jp/, Tel: 81-45-503-9222,
Fax: 81-45-503-9216)
COMMENT
cDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.
Please visit our web site for further details.
URL: http://genome.gsc.riken.jp/
URL: http://fantom.gsc.riken.jp/
FEATURES
Location/Qualifiers
1..4261
/organism="Mus musculus"
/mol_type="mRNA"
/strain="CS7BL/6J"
/db_xref="FANTOM,DB:D130043C18"
/db_xref="taxon:10090"
/clone="D130043C18"
/tissue_type="spinal ganglion"
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Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTTAGCT 21
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Db 1497 CAGTCATATGCAGGCTTAGCT 1477

RESULT 7
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Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCTAGC 20
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pb 314 CAGTANCATGCAGGTCTAGC 333

[illegible]

ORGANISM

Zea mays

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD clade; Panicoideae; Andropogoneae; Zea.

AUTHORS
Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,
Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T.,
Citek, R.W., Nunberg, A., Robbins, D. and Lakey, N.

TITLE Consortium for Maize Genomics
JOURNAL Unpublished (2002)
COMMENT Contact: Cathy Whitelaw
TIGR

TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA

Tel: 301-838-5843

Fax: 301-838-0208

Email: whitelaw@tigr.org

Seq primer: TR

Class: sheared ends.

FEATURES	Location/Qualifiers
source	1. .772

ORIGIN

Query Match 82.9%; Score 17.4; DB 8; Length 772;
Best Local Similarity 94.7%; Pred. No. 1.2e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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DEFINITION	OGSCS93TC ZM 0.7 1.5 KB Zea mays genomic clone ZMMBm0832B05, genomic survey sequence.			GSS 22-AUG-2003

ORGANISM	Zea mays
Eukaryota;	Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta;	Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoidae; Andropogoneae; Zea.	

REFERENCE
AUTHORS
1 (bases 1 to 772)
Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,
Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T.,
Citek, R.W., Nunberg, A., Robbins, D. and Lakey, N.

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCTAG 19

Db 384 CATTGACATGCAGGTCTAG 366

RESULT 13

B1548151/c

LOCUS

DEFINITION

603189492F1 NIH_MGC_95 Homo sapiens cDNA clone IMAGE:5260847 5',

mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

1 (bases 1 to 816)

NIH-MGC http://mgc.nci.nih.gov/.

Unpublished (1999)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-remail.nih.gov

Tissue Procurement: Miklos Palkovits, M.D., Ph.D.

cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki

Toshiyuki and Pietro Carninci (RIKEN)

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

http://image.llnl.gov

Plate: LLAM11657 row: 9 column: 24

High quality sequence stop: 740.

FEATURES

source

1..816

Location/Qualifiers

/organism="Homo sapiens"

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/clone="IMAGE:5260847"

/tissue_type="hippocampus"

/lab_host="DH10B"

/clone_lib="NIH MGC 95"

/notes="Organ: brain; Vector: pBluescriptR (modified

pBluescript KS+); Site 1: BamHI; Site 2: SalI-XhoI

(gtcgag); Oligo-dT primed using primer

5'-TTTTTTTTTTTTTTVN-3', size-selected for average

insert size 2.5 kb and normalized to R0T 5. This is a

primary library enriched for full-length clones and

constructed using the Cap-trapper method (Carninci, in

preparation). Library constructed by M. Brownstein

(NIH/NHGRI, National Institutes of Health). Note: this

is a NIH_MGC Library."

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

1 (bases 1 to 816)

NIH-MGC http://mgc.nci.nih.gov/.

Unpublished (1999)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-remail.nih.gov

Tissue Procurement: Miklos Palkovits, M.D., Ph.D.

cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki

Toshiyuki and Pietro Carninci (RIKEN)

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Incyte Genomics, Inc.

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

1 (bases 1 to 817)

Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,

Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T.,

Citek, R.W., Nunberg, A., Robbins, D. and Lakey, N.

Consortium for Maize Genomics

Unpublished (2002)

Contact: Cathy Whitelaw

TIGR

9712 Medical Center Drive, Rockville, MD 20850, USA

Tel: 301-838-5843

Fax: 301-838-0208

Email: whitelaw@tigr.org

Seq primer: TF

Class: sheared ends.

Location/Qualifiers

1..817

/organism="Zea mays"

/mol_type="genomic DNA"

/strain="B73"

/db_xref="taxon:4577"

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methylation filtered genomic DNA library"

ORIGIN

Query Match

Best Local Similarity

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCTAG 19

Db 488 CATTGACATGCAGGTCTAG 506

RESULT 15

CG211861/c

LOCUS

DEFINITION

CG211861

CG211861

CG211861.1

CG211861.1

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Zea mays

Zea mays

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD

clade; Panicoideae; Andropogoneae; Zea.

1 (bases 1 to 817)

Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,

Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T.,

Citek, R.W., Nunberg, A., Robbins, D. and Lakey, N.

Consortium for Maize Genomics

Unpublished (2002)

Contact: Cathy Whitelaw

TIGR

9712 Medical Center Drive, Rockville, MD 20850, USA

Tel: 301-838-5843

Fax: 301-838-0208

Email: whitelaw@tigr.org

Seq primer: TF

Class: sheared ends.

Location/Qualifiers

1..817

/organism="Zea mays"

/mol_type="genomic DNA"

/strain="B73"

/db_xref="taxon:4577"

/clone_lib="ZM 0.7 1.5 KB"

/clone="ZMMBMA0368M23"

/note="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb

methylation filtered genomic DNA library"

ORIGIN

Query Match

Best Local Similarity

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCTAG 19

Db 488 CATTGACATGCAGGTCTAG 506

RESULT 15

CG211861/c

LOCUS

DEFINITION

CG211861

CG211861

CG211861.1

CG211861.1

CG211861.1

CG211861.1

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CG211861.1

Zea mays

Zea mays

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD

clade; Panicoideae; Andropogoneae; Zea.

1 (bases 1 to 817)

Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,

Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T.,

Citek, R.W., Nunberg, A., Robbins, D. and Lakey, N.

Consortium for Maize Genomics

Unpublished (2002)

Contact: Cathy Whitelaw

TIGR

9712 Medical Center Drive, Rockville, MD 20850, USA

Tel: 301-838-5843

Fax: 301-838-0208

Email: whitelaw@tigr.org

Seq primer: TF

Class: sheared ends.

Location/Qualifiers

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/organism="Zea mays"

/mol_type="genomic DNA"

/strain="B73"

/db_xref="taxon:4577"

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/note="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb

methylation filtered genomic DNA library"

ORIGIN

Query Match

Best Local Similarity

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCTAG 19

Db 488 CATTGACATGCAGGTCTAG 506

RESULT 15

CG211861/c

LOCUS

DEFINITION

CG211861

CG211861

CG211861.1

CG211861.1

CG211861.1

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ORIGIN
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Best Local Similarity 94.7%; Pred. No. 1.2e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 157 CATTGACATGCAGGTCCTAG 139

RESULT 16
BY609340
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

REFERENCE
AUTHORS

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 382)
Okazaki,Y., Furuno,M., Kasukawa,T., Adachi,J., Bono,H., Kondo,S.,
Nikaido,I., Osato,N., Saito,R., Suzuki,H., Yamanaka,I.,
Kiyosawa,H., Yagi,K., Tomaru,Y., Hasegawa,Y., Nogami,A.,
Hume,D.A., Quackenbush,J., Schriml,L.M., Kanapin,A., Bult,C.,
Batalov,S., Beisel,K.W., Blake,J.A., Bradt,D., Brusci,V.,
Chochia,C., Corbani,L.E., Cousins,S., Dalla,E., Dragani,T.A.,
Fletcher,C.F., Forrest,A., Frazer,K.S., Gaasterland,T.,
Cariboldi,M., Gissi,C., Godzik,A., Gough,J., Grimmond,S.,
Gustincich,S., Hirokawa,N., Jackson,I.J., Jarvis,E.D., Kanai,A.,
Kawaji,H., Kawasawa,Y., Kedzierski,R.M., King,B.L., Konagaya,A.,
Kurochkin,I.V., Lee,Y., Lenhard,B., Lyons,P.A., Maglott,D.R.,
Maltais,L., Marchionni,L., McKenzie,L., Miki,H., Nagashima,T.,
Numata,K., Okido,T., Pavan,W.J., Pertea,G., Pesole,G.,
Petrovsky,N., Pillai,R., Pontius,J.U., Qi,D., Ramachandran,S.,
Ravasi,T., Reed,D.J., Reid,J., Ring,B.Z., Ringwald,M.,
Sandelin,A., Schneider,C., Semple,C.A., Setou,M., Shimada,K.,
Sultana,R., Takenaka,Y., Taylor,M.S., Teasdale,R.D., Tomita,M.,
Verardo,R., Wagner,L., Wahlestedt,C., Wang,Y., Watanabe,Y.,
Wells,C., Wilming,L.G., Wynnshaw-Boris,A., Yanagisawa,M., Yang,I.,
Yang,L., Yuan,Z., Zavolan,M., Zhu,Y., Zimmer,A., Carninci,P.,
Hayatsu,N., Hirozane-Kishikawa,T., Konno,H., Nakamura,M.,
Sakazume,N., Sato,K., Shiraki,T., Waki,K., Kawai,J., Aizawa,K.,
Arakawa,T., Fukuda,S., Hara,A., Hashizume,W., Imotani,K., Ishii,Y.,
Itoh,M., Kagawa,I., Miyazaki,A., Sakai,K., Sasaki,D., Shibata,K.,
Shinagawa,A., Yasunishi,A., Yoshino,M., Waterston,R., Lander,E.S.,
Rogers,J., Birney,E. and Hayashizaki,Y.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)
22354683
PUBMED
12466851
COMMENT
Contact: Yoshhide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-resesgsc.riken.jp, URL:http://genome.gsc.riken.jp/
Aizawa,K., Akimura,T., Arakawa,T., Carninci,P., Fukuda,S.,
Hirozane,T., Imotani,K., Ishii,Y., Itoh,M., Kawai,J., Konno,H.,
Miyazaki,A., Murata,M., Nakamura,M., Nomura,K., Numazaki,R.,
Ohno,M., Sakai,K., Sakazume,N., Sasaki,D., Sato,K., Shibata,K.,

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Shiraki,T., Tagami,M., Waki,K., Watahiki,A., Muramatsu,M. and
Hayashizaki,Y. Direct Submission
Computational Analysis of Full-length Mouse cDNAs Compared with
Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new
genes. Genome Res. 10 (10), 1617-1630 (2000)
RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multicapillary sequencer. Genome Res.
10 (11), 1757-1771 (2000)
Computer-based methods for the mouse full-length cDNA
encyclopedia: real-time sequence clustering for construction of a
nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
cDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.
Tissues were provided by Michela Fagioli and Takao K. Henssch (
Laboratory for Neuronal Circuit Development Brain Science Institute
RIKEN 2-1 Hirosawa,Wako-shi,Saitama 351-0198 Japan ) whose
assistance we gratefully acknowledge.
Please visit our web site (http://genome.gsc.riken.go.jp) for
further details.
Location/Qualifiers
1..382
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/mol_type="mRNA"
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QY 2 AGTGACATGCAGGTCCTA 18
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Db 28 AGTGACATGCAGGTCCTA 44

RESULT 17
BY649695
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

REFERENCE
AUTHORS

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 408)
Okazaki,Y., Furuno,M., Kasukawa,T., Adachi,J., Bono,H., Kondo,S.,
Nikaido,I., Osato,N., Saito,R., Suzuki,H., Yamanaka,I.,
Kiyosawa,H., Yagi,K., Tomaru,Y., Hasegawa,Y., Nogami,A.,
Schonbach,C., Gojohori,T., Baldarelli,R., Hill,D.P., Bult,C.,
Hume,D.A., Quackenbush,J., Schriml,L.M., Kanapin,A., Matsuda,H.,
Batalov,S., Beisel,K.W., Blake,J.A., Bradt,D., Brusci,V.,
Chochia,C., Corbani,L.E., Cousins,S., Dalla,E., Dragani,T.A.,
Fletcher,C.F., Forrest,A., Frazer,K.S., Gaasterland,T.,
Gustincich,S., Hirokawa,N., Jackson,I.J., Jarvis,E.D., Kanai,A.,
Kawaji,H., Kawasawa,Y., Kedziereki,R.M., King,B.L., Konagaya,A.,
Kurochkin,I.V., Lee,Y., Lenhard,B., Lyons,P.A., Maglott,D.R.,
Maltais,L., Marchionni,L., McKenzie,L., Miki,H., Nagashima,T.,
Numata,K., Okido,T., Pavan,W.J., Pertea,G., Pesole,G.,
Petrovsky,N., Pillai,R., Pontius,J.U., Qi,D., Ramachandran,S.,
Ravasi,T., Reed,D.J., Reid,J., Ring,B.Z., Ringwald,M.,

```

Sandelin, A., Schneider, C., Sempke, C.A., Setou, M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, M.S., Teasdale, R.D., Tomita, M., Uetano, R., Wagner, L., Wahlstedt, C., Wang, Y., Watanabe, Y., Wells, C., Wilming, L.G., Wyshaw-Boris, A., Yanagisawa, M., Yang, I., Yang, L., Yuan, Z., Zavalan, M., Zhu, Y., Zimmer, A., Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura, M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K., Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K., Shinagawa, A., Yasunishi, A., Yoshino, M., Watanabe, R., Lander, E.S., Rogers, J., Birney, E. and Hayashizaki, Y.

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Nature 420, 563-573 (2002)
22354683

Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Susehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216

Email: genome-res@gsc.riken.jp, URL: <http://genome.gsc.riken.jp/>
Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hirozane, T., Imotani, K., Ishii, Y., Itoh, M., Kawai, J., Konno, H., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Waki, K., Watahiki, A., Muramatsu, M. and Hayashizaki, Y. Direct Submission

Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)

Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)

RIKEN integrated sequence analysis (RISA) system-384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Tissues were provided by Michela Pagiolini and Takao K. Hensch (Laboratory for Neuronal Circuit Development Brain Science Institute RIKEN 2-1 Hiroswa, Wako-shi, Saitama 351-0198 Japan) whose assistance we gratefully acknowledge.

Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

FEATURES
source

BEST LOCAL SIMILARITY 100.0%; FREQ. NO. 1.8E+03;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 TGACATGCAGGTCTAGC 20
Db 444 TGACATGCAGGTCTAGC 428

ORIGIN

Query Match	81.0%	Score 17;	DB 6;	Length 408;
Best Local Similarity	100.0%;	Pred. No. 1.7e+03;		
Matches 17;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

Qy 2 AGTGACATGCAGGTCTA 18
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Db 56 AGTGACATGCAGGTCTA 72

RESULT 18

CA124343/C
LOCUS
DEFINITION

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

REFERENCE

REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT
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FEATURES SOURCE

ORIGIN

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RESULT 19
AZ069535

LOCUS
DEFINITION
ACCESSION

VERSION
KEYWORDS
SOURCE
ORGANISM

REFERENCE
AUTHORS

Akinret, B., Levins, M., Mcgann, S., Tsagaye, G., Geer, K., Krol, M., de Jong, P. and Fraser, C.M. BAC End Sequences from Library RPCI-23 Unpublished (1999)
 Other GSSs: RPCI-23-435G22-TV
 Contact: Shaying Zhao
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: szhao@tigr.org
 Clones are derived from the mouse BAC library RPCI-23. For BAC library availability, please contact Pieter de Jong (pieterdejong.med.buffalo.edu). Clones may be purchased from BPAC Resources (<http://bacpac.med.buffalo.edu/orderingframe.htm>) or from Resea ch Genetics (info@resgen.com). BAC end page: http://www.tigr.org/cdb/bac/bac_ends/mouse/bac_end_intro.html
 Plate: 435 row: G column: 22
 Seq primer: SP6
 Class: BAC ends.

FEATURES

source

Location/Qualifiers

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1..633
/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="RPCI-23-435G22"
/sex="Female"
/lab_host="DH10B"
/clone_lib="RPCI-23"
/note="Organ: Kidney/Brain; Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI; Female C57BL/6J mouse kidney and/or brain genomic DNA was isolated and partially digested with a combination of EcoRI and EcoRI Methylase. Size selected DNA was cloned into the pBACe3.6 vector at the EcoRI sites. The ligation products were transformed into DH10B electrocompetent cells (BRL Life Technologies)."
```

ORIGIN

```
Query Match      81.0%; Score 17; DB 8; Length 633;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY 1 CAGTGACATGCAGGTCT 17

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Db 122 CAGTGACATGCAGGTCT 138

RESULT 20

BB650662/c

LOCUS

```
DEFINITION      BB650662 RIKEN full-length enriched, 0 day neonate cerebellum Mus
musculus cDNA clone C230020D16 5', mRNA sequence.
```

ACCESSION

BB650662

VERSION

EST.

KEYWORDS

Mus musculus (house mouse)

ORGANISM

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 642)

Arakawa, T., Carninci, P., Fukuda, S., Furuno, M., Hangsaki, T.,

Hara, A., Hiramoto, K., Hori, F., Ishii, Y., Ito, M., Kawai, J.,

Konno, H., Kouda, M., Koya, S., Matsuyama, T., Miyazaki, A., Nomura, K.,

Ohno, M., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K.,

Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T.,

Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F.,

Takeda, Y., Tanaka, T., Toyota, T., Muramatsu, M. and Hayashizaki, Y.

RIKEN Mouse ESTs (Arakawa, T., et al. 2001)

Unpublished (2001)

Contact: Yoshihide Hayashizaki

Laboratory for Genome Exploration Research Group, RIKEN Genomic

Sciences Center (GSC), Yokohama Institute
 The Institute of Physical and Chemical Research (RIKEN)
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
 Tel: 81-45-503-9222
 Fax: 81-45-503-9216

Email: genome-res@gsc.riken.jp, URL: <http://genome.gsc.riken.jp/>
 Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
 Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
 Normalization and subtraction of cap-trapper-selected cDNAs to
 prepare full-length cDNA libraries for rapid discovery of new
 genes. Genome Res. 10 (10), 1617-1630 (2000)

wagi, K., Fujiwaki, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,
 Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T.,
 Matsumura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A.
 and Hayashizaki, Y.

RIKEN integrated sequence analysis (RISA) system--384-format
 sequencing pipeline with 384 multicapillary sequencer. Genome Res.
 10 (11), 1757-1771 (2000)

Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P.,
 Sugahara, Y. and Hayashizaki, Y.

Computer-based methods for the mouse full-length cDNA
 encyclopedia: real-time sequence clustering for construction of a
 nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

Kondo, S., Shinagawa, A., Saito, T., Kiyosawa, H., Yamanaka, I.,
 Aizawa, K., Fukuda, S., Hara, A., Itoh, M., Kawai, J., Shibata, K. and
 Hayashizaki, Y.

Computational Analysis of Full-Length Mouse cDNAs Compared with
 Human Genome Sequences. Mamm. Genome. 12, 673-677 (2001)
 Please visit our web site (<http://genome.gsc.riken.go.jp>) for
 further details.

e mouse tissues.

FEATURES

source

Location/Qualifiers

1..642

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="C230020D16"

/tissue_type="cerebellum"

/dev_stage="0 day neonate"

/lab_host="DH10B"

/clone_lib="RIKEN full-length enriched, 0 day neonate

cerebellum"

/note="Site 1: SalI; Site 2: BamHI; cDNA library was

prepared and sequenced in Mouse Genome Encyclopedia

Project of Genome Exploration Research Group in Riken

Genomic Sciences Center and Genome Science Laboratory in

RIKEN. Division of Experimental Animal Research in Riken

contributed to prepare mouse tissues. 1st strand cDNA was

primed with a primer [5'

GAGAGAGAGAGATCCAGAGCTCTTTTTTTTTTTTNN 3'], cDNA was

prepared by using trehalose thermo-activated reverse

transcriptase and subsequently enriched for full-length by

cap-trapper. cDNA went through one round of normalization

to Rot = 20.0 and subtraction to Rot = 479.0. Second

strand cDNA was prepared with the primer adapter of

sequence [5' GAGAGAGATTCGAGTTAATAATATCCGCCCCCCCCC

3']. cDNA was cleaved with XhoI and BamHI. Vector: a

modified phluescript KS(+) after bulk excision from Lambda

FLC I."

ORIGIN

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Query Match      81.0%; Score 17; DB 2; Length 642;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY 1 CAGTGACATGCAGGTCT 17

|||||

Db 27 CAGTGACATGCAGGTCT 11

|||||

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|||||

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|||||

RESULT 21

BB293162/c

LOCUS

BB293162

655 bp

mRNA

linear

EST 24-OCT-2001

DEFINITION BB293162 RIKEN full-length enriched, 9.5 days embryo parthenogenote Mus musculus cDNA FLJ10755 fis, clone NT2RP3004569, weakly similar to sapiens cDNA FLJ10755 fis, clone NT2RP3004569, weakly similar to ANKYRIN, BRAIN VARIANT 1, mRNA sequence.

ACCESSION BB293162

VERSION BB293162.2

KEYWORDS GI:16401650

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

REFERENCE Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 655)

AUTHORS Arakawa,T., Carninci,P., Fukuda,S., Furuno,M., Hanagaki,T., Hara,A., Hiramoto,K., Hori,F., Ishii,Y., Ito,M., Kawai,J., Konno,H., Kouda,M., Koya,S., Matsuyama,T., Miyazaki,A., Nomura,K., Ohno,M., Okazaki,Y., Okido,T., Saito,R., Sakai,C., Sakai,K., Sano,H., Sasaki,D., Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H., Tagami,M., Tagawa,A., Takahashi,F., Takeda,Y., Tanaka,T., Toya,T., Muramatsu,M. and Hayashizaki,Y. RIKEN Mouse ESTs (Arakawa,T., et al. 2001) Unpublished (2001)

TITLE On Jul 10, 2000 this sequence version replaced gi:8993654.

JOURNAL Contact: Yoshihide Hayashizaki

COMMENT Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute The Institute of Physical and Chemical Research (RIKEN) 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222 Fax: 81-45-503-9216 Email: genome-resgsc.riken.jp, URL: http://genome.gsc.riken.jp/ Konno,H., Fukunishi,Y., Shibata,K., Itoh,M., Carninci,P., Sugahara,Y. and Hayashizaki,Y. Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001) Kondo,S., Shinagawa,A., Saito,T., Kiyosawa,H., Yamanaoka,I., Aizawa,K., Fukuda,S., Hara,A., Itoh,M., Kawai,J., Shibata,K. and Hayashizaki,Y. Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001) Please visit our web site (http://genome.gsc.riken.go.jp/) for further details. cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in RIKEN Genomic Sciences Center and Genome Science Laboratory in RIKEN Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

FEATURES Location/Qualifiers

1..655

 /organism="Mus musculus"

 /mol_type="mRNA"

 /db_xref="taxon:10090"

 /clone="BI30017A20"

 /tissue_type="parthenogenote"

 /dev_stage="9.5 days embryo"

 /lab_host="DH10B"

 /clone_lib="RIKEN full-length enriched, 9.5 days embryo parthenogenote"

 /notes="Site 1: Sali; Site 2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken"

ORIGIN

Query Match 81.0%; Score 17; DB 2; Length 655;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCT 17
 |||||
 Db 145 CAGTGACATGCAGGTCT 129

RESULT 22
 AG457137 768 bp DNA linear GSS 04-JUN-2004
 LOCUS Mus musculus molossinus DNA, clone:MSMg01-343p07.TJ, genomic survey sequence.

DEFINITION AG457137
 VERSION AG457137.1 GI:48148651
 KEYWORDS GSS.
 SOURCE Mus musculus molossinus
 ORGANISM Mus musculus molossinus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1
 Hattori,M., Toyoda,A., Noguchi,H., Kojima,T. and Sakaki,Y.
 BAC end Sequences of Library MSMg01
 Unpublished
 2 (bases 1 to 768)
 Direct Submission
 Submitted (17-NOV-2003) Masahira Hattori, The Institute of Physical and Chemical Research (RIKEN), Genomic Sciences Center (GSC); 1-7-22 Suehiro-cho,Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail:hattori@gsc.riken.jp, URL:http://hgp.gsc.riken.go.jp/, Tel:81-45-503-9111, Fax:81-45-503-9170)
 Clones are derived from the mouse BAC library MSMg01. For BAC library availability, please contact Kuniya Abe (abe@rtc.riken.jp).
 Tsukuba Institute, Bio Resource Center,
 The Institute of Physical and Chemical Research (RIKEN) 3-1-1
 Koyada, Tsukuba, 305-0074 Japan
 phone: 81-298-36-9189, fax: 81-298-36-9199
 e-mail: abe@rtc.riken.jp

COMMENT Sequencing : TJ
 PRIMERS
 LIBRARY
 Vector : pBACe3.6
 R.Site 1 : EcoRI
 R.Site 2 : EcoRI.

FEATURES Location/Qualifiers

1..768

 /organism="Mus musculus molossinus"

 /mol_type="genomic DNA"

 /sub_species="molossinus"

 /db_xref="taxon:57486"

 /clone="MSMg01-343p07.TJ"

 /sex="male"

 /tissue_type="mixture of kidney and spleen"

 /clone_lib="MSMg01 Mouse Male BAC Library"

ORIGIN

```

Query Match      81.0%; Score 17; DB 9; Length 768;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCT 17
    |||||
Db 191 CAGTGACATGCAGGTCT 207

RESULT 23
AG557182      781 bp DNA linear GSS 05-JUN-2004
LOCUS Mus musculus molossinus DNA, Clone:MSMg01-475P07.T7, genomic survey
DEFINITION
ACCESSION AG557182
VERSION AG557182.1 GI:48317880
KEYWORDS GSS.
SOURCE Mus musculus molossinus
ORGANISM Mus musculus molossinus
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Euthera; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
Hattori,M., Toyoda,A., Noguchi,H., Kojima,T. and Sakaki,Y.
BAC end Sequences of Library MSMg01
TITLE Unpublished
JOURNAL
REFERENCE Hattori,M., Toyoda,A., Noguchi,H., Kojima,T. and Sakaki,Y.
AUTHORS BAC end Sequences of Library MSMg01
TITLE Unpublished
JOURNAL
REFERENCE Hattori,M., Toyoda,A., Noguchi,H., Kojima,T. and Sakaki,Y.
AUTHORS Direct Submission
TITLE Direct Submission
JOURNAL
COMMENT 17-NOV-2003 Masahira Hattori, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
Submitted (17-NOV-2003) Masahira Hattori, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
1-7-22 Suehiro-chou,Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
(E-mail:hattori@gsc.riken.jp, URL:http://hgpc.gsc.riken.go.jp/,
Tel:81-45-503-9111, Fax:81-45-503-9170)
Clones are derived from the mouse BAC library MSMg01. For BAC
library availability, please contact Kuniya Abe (abe@tc.riken.jp).
Teskuba Institute, Bio Resource Center.
The Institute of Physical and Chemical Research (RIKEN) 3-1-1
Koyadai, Tsukuba, 305-0074 Japan
phone: 81-298-36-9189, fax: 81-298-36-9199
e-mail: abe@tc.riken.jp
PRIMERS
Sequencing : T7
LIBRARY
Vector : pBAC3.6
R.Site 1 : EcoRI
R.Site 2 : EcoRI.
FEATURES
Location/Qualifiers
source
1..781
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/mol_type="genomic DNA"
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/clone="MSMg01-475P07.T7"
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ORIGIN
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Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCT 17
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Db 187 CAGTGACATGCAGGTCT 203

RESULT 24
CF411086      949 bp mRNA linear EST 02-SEP-2003
LOCUS CH3#071_D10MF Canine heart normalized cDNA Library in pBluescript
DEFINITION
Canis familiaris cDNA clone CH3#071_D10 5', mRNA sequence.
CF411086
ACCESSION

Query Match      81.0%; Score 17; DB 7; Length 949;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCT 17
    |||||
Db 299 CAGTGACATGCAGGTCT 315

RESULT 25
AK044974/c     2445 bp mRNA linear HTC 03-APR-2004
LOCUS Mus musculus 9.5 days embryo parthenogenote cDNA, RIKEN full-length
DEFINITION enriched library, clone.B130017A20 product:synuclein, alpha
interacting protein (synphilin), full insert sequence.
AK044974
ACCESSION AK044974.1 GI:26336964
VERSION AK044974.1
KEYWORDS HTC; CAP trapper.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Euthera; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
Carninci,P. and Hayashizaki,Y.
TITLE High-efficiency full-length cDNA cloning
JOURNAL Meth. Enzymol. 303, 19-44 (1999)
MEDLINE 99279253
PUBMED 10349636
REFERENCE Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K.,
AUTHORS Itoh,M., Konno,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.

```

TITLE Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new genes
JOURNAL Genome Res. 10 (10), 1617-1630 (2000)
MEDLINE 20499374
PUBMED 11042159
REFERENCE 3

AUTHORS Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
Konno, H., Akiyama, J., Nishi, K., Kitsuunai, T., Tashiro, H., Itoh, M.,
Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A.,
Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K.,
Fujiwara, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M.,
Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J.,
Okazaki, Y., Murakami, M., Inoue, Y., Kira, A., and Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multicapillary sequencer
Genome Res. 10 (11), 1757-1771 (2000)

TITLE The RIKEN Genome Exploration Research Group Phase II Team and the
FANTOM Consortium.
JOURNAL Functional annotation of a full-length mouse cDNA collection
MEDLINE Nature 409, 685-690 (2001)
PUBMED 11076861
REFERENCE 4

AUTHORS The RIKEN Consortium and the RIKEN Genome Exploration Research
Group Phase I & II Team.
TITLE Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
JOURNAL Nature 420, 563-573 (2002)
REFERENCE 6 (bases 1 to 2445)

AUTHORS Adachi, J., Aizawa, K., Akimura, T., Arawaka, T., Bono, H., Carninci, P.,
Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W.,
Hayashida, K., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T.,
Hori, P., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kasukawa, T.,
Kato, H., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Kouda, M.,
Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, M.,
Nakamura, M., Nishi, K., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N.,
Okazaki, Y., Saito, R., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N.,
Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T.,
Sogabe, Y., Tagami, M., Tagawa, A., Takahashi, F., Takaku-Akahira, S.,
Takeda, Y., Tanaka, T., Tomaru, A., Toyota, T., Yasunishi, A.,
Muramatsu, M., and Hayashizaki, Y.

TITLE Direct Submission
JOURNAL Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of
Physical and Chemical Research (RIKEN), Laboratory for Genome
Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
RIKEN Yokohama Institute, 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
Kanagawa 230-0045, Japan (E-mail: genome-res@gsc.riken.jp,
URL: http://genome.gsc.riken.jp/, Tel: 81-45-503-9222,
Fax: 81-45-503-9216)

COMMENT cDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.

Tissues were provided by Dr. Tomohiro Kono (Department of Animal
Science, Tokyo University of Agriculture, 1737 Hunko Atsugi City,
Kanagawa Prefecture, Japan) whose assistance we gratefully
acknowledge.

Please visit our web site for further details.
URL: http://genome.gsc.riken.jp/
URL: http://fantom.gsc.riken.jp/.

FEATURES Location/Qualifiers
source 1..2445
organism="Mus musculus"
mol_type="mRNA"
strain="C57BL/6J"
db_xref="FANTOM DB: B130017A20"
db_xref="taxon: 10090"
clone="B130017A20"
tissue_type="parthenogenote"
clone_lib="RIKEN full-length enriched mouse cDNA library"
dev_stage="9.5 days embryo"

CDS

385..>2445
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(MGD|MGJ:1915097, GB|NM_026408, evidence: BLASTN, 99%,
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/protein_id="BAC32166.1"
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EYKGETDQGPQPELSPEDGVGGLPGKGSFQALGSEHYDLDMDLIDVPIYKS
SOOLAPLTVTSEKRIILGCTTINGLSAKTPIASTENSTPNNTPCVLSPVKSPLHR
KAPTRLDQHLSTEDSSPAGKCGPAYESNHSKDFLNKVFSPHKKIKSGPDP
CLAAONLDKIHDENGNLLHIAASKGAECLOHLTSLMGEDCLNENTEQLTPAGLAI
HQAOLECVRMVYSETEAIAELSCDFPSLIHYAGCYGOEKILLWLOHMOEGISLD
EVDRENSAVHVASOHGYLGCITLVEYGANTVMQNHAGEKSPQSQAERHGHILCSRYL
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NSEDP"

ORIGIN

Query Match 81.0%; Score 17; DB 3; Length 2445;
Best Local Similarity 100.0%; Pred. No. 2.3e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CAGTGACATGCAGGTCT 17
Db 1935 CAGTGACATGCAGGTCT 1919

Search completed: September 6, 2005, 21:56:00
Job time : 1506.84 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 16:01:23 ; Search time 189.656 Seconds
(without alignments)
655.473 Million cell updates/sec

Title: US-10-729-421-40

Perfect score: 21

Sequence: 1 cagtgcacatcgaggtcttagct 21

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 100 summaries

Database : N_Geneseq_16Dec04:*

1: Geneseqn1980s:*

2: Geneseqn1990s:*

3: Geneseqn2000s:*

4: Geneseqn2001as:*

5: Geneseqn2001bs:*

6: Geneseqn2002as:*

7: Geneseqn2002bs:*

8: Geneseqn2003as:*

9: Geneseqn2003bs:*

10: Geneseqn2003cs:*

11: Geneseqn2003ds:*

12: Geneseqn2004as:*

13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	21	100.0	21	8	Acc80536 Exemplary
2	21	100.0	21	8	Acc80538 Internal
3	21	100.0	21	8	Acc80537 Internal
4	21	100.0	21	9	Abz59632 Parvoviru
5	21	100.0	21	12	Adi53796 HAV inter
6	21	100.0	23	12	Adi53797 HAV inter
7	21	100.0	23	12	Adq30670 West Nile
8	21	100.0	25	12	Adq30671 West Nile
9	21	100.0	681	9	Abz59634 Exemplary
10	21	100.0	727	12	Adi53795 HAV inter
11	21	100.0	967	12	Adq30647 West Nile
12	21	100.0	1696	8	Acc80539 Internal
13	17.4	82.9	5446	10	Adi71204 Novel hum
14	17.4	82.9	28482	8	Abz73855 Secreted
15	17.4	82.9	28482	8	Adi4262 Human sec
16	17.4	82.9	32681	8	Abz74517 Secreted
17	17.4	82.9	32681	8	Abz73854 Secreted
18	17.4	82.9	32681	8	Adi98915 Human sec
19	17.4	82.9	32681	8	Adi44261 Human sec
20	17.4	82.9	32681	8	Adi44519 Human sec

C 21	17.4	82.9	32681	10	ADC20949	Adc20949 Human sec
C 22	17.4	82.9	32681	10	ABZ68053	Abz68053 Human sec
C 23	16.8	80.0	320	8	ABX46275	Abx46275 Bovine ES
C 24	16.8	80.0	36221	4	AAS00624	Aas00624 Human dea
C 25	16.4	78.1	72705	11	ACN45158	Acn45158 Human gen
C 26	16.4	78.1	110000	8	ABX16390_5	Continuation 16 of
C 27	16.4	78.1	117382	11	ACN44804	Acn44804 Mouse gen
C 28	16.4	78.1	340449	8	AAL52198	Aal52198 Human sec
C 29	16.2	77.1	201	13	ADS40801	Adi40801 Human aut
C 30	16.2	77.1	201	13	ADS39530	Adi39530 Human aut
C 31	16.2	77.1	394	4	AAL82251	Aal82251 Human pol
C 32	16.2	77.1	473	9	ACH36560	Ach36560 Human end
C 33	16.2	77.1	497	10	ADB56505	Adi56505 Toxicity-
C 34	16.2	77.1	1487	10	ADC71327	Adc71327 Human col
C 35	16.2	77.1	2048	2	AAQ85985	Aaq85985 Zea maye
C 36	16.2	77.1	2174	10	ADE62245	Ade62245 Rat gene
C 37	16.2	77.1	2238	5	AAS76384	Aas76384 DNA encod
C 38	16.2	77.1	2569	13	ADR08803	Adr08803 Full leng
C 39	16.2	77.1	2721	8	AAL53547	Aal53547 cDNA of h
C 40	16.2	77.1	4445	6	ABAO1096	Abao1096 Brevibact
C 41	16.2	77.1	9048	4	AAC90812	Aac90812 B. lactof
C 42	16.2	77.1	10500	4	AAL05334	Aal05334 Human rep
C 43	16.2	77.1	10500	4	ABL98203	AbL98203 Human tes
C 44	16.2	77.1	14902	13	ADS36489	Adi36489 Human aut
C 45	16.2	77.1	15515	8	AAL53548	Aal53548 Genomic D
C 46	16.2	77.1	55827	8	ACA60949	AcA60949 DNA encod
C 47	16.2	77.1	58337	13	ADS36454	Adi36454 Human aut
C 48	16.2	77.1	64423	13	ADS36462	Adi36462 Human aut
C 49	16.2	77.1	70372	6	AAL53466	Aal53466 Rab-like
C 50	16.2	77.1	90442	9	ADA03077	Ada03077 Mouse mCG
C 51	16.2	77.1	90442	9	ADA66361	Ada66361 Mouse mCG
C 52	16.2	77.1	90442	10	ADB72815	Adi72815 Mouse mCG
C 53	16.2	77.1	90442	10	ADC26997	Adc26997 Mouse car
C 54	16.2	77.1	90442	11	ADL27155	AdL27155 Mouse gen
C 55	16.2	77.1	143306	6	ABK49586	Abk49586 Human tra
C 56	16.2	77.1	349980	5	AAH68529	Aah68529 C glutami
C 57	15.8	75.2	171	2	AAV89101	Aav89101 EST clone
C 58	15.8	75.2	279	2	AAV87960	Aav87960 EST clone
C 59	15.8	75.2	297	9	ADB08791	Adi08791 Alloiococ
C 60	15.8	75.2	297	9	ADB08789	Adi08789 Alloiococ
C 61	15.8	75.2	497	5	AAS88145	Aas88145 DNA encod
C 62	15.8	75.2	497	5	AAS80088	Aas80088 DNA encod
C 63	15.8	75.2	587	4	AAH07183	Aah07183 Human cDN
C 64	15.8	75.2	748	4	AAH03978	Aah03978 Human cDN
C 65	15.8	75.2	882	10	ADC08897	Adc08897 Rice DNA
C 66	15.8	75.2	1614	9	ADB08797	Adi08797 Alloiococ
C 67	15.8	75.2	1670	4	AAH17173	Aah17173 Human cDN
C 68	15.8	75.2	1777	4	AAH16391	Aah16391 Human cDN
C 69	15.8	75.2	1995	6	ABL88382	AbL88382 Pain regu
C 70	15.8	75.2	2378	5	AAS64699	Aas64699 DNA encod
C 71	15.8	75.2	2379	5	AAS67041	Aas67041 DNA encod
C 72	15.8	75.2	4015	12	ADJ34728	Adj34728 Rat 2'-5'
C 73	15.8	75.2	4708	12	ADJ34707	Adj34707 Mouse 2'-
C 74	15.8	75.2	31236	9	ADA02900	Ada02900 Human PTP
C 75	15.8	75.2	31236	10	ADB72638	Adi72638 Human PTP
C 76	15.8	75.2	31236	10	ADC85379	Adc85379 Mouse PTP
C 77	15.8	75.2	31236	12	ADM74495	Adm74495 Human car
C 78	15.8	75.2	31718	4	AAK90359	Aak90359 Human dig
C 79	15.8	75.2	31718	4	AAK90360	Aak90360 Human dig
C 80	15.8	75.2	31718	4	AAK73104	Aak73104 Human imm
C 81	15.8	75.2	31718	4	AAK87573	Aak87573 Human imm
C 82	15.8	75.2	31718	4	AAK73120	Aak73120 Human imm
C 83	15.8	75.2	31718	4	AAK87442	Aak87442 Human imm
C 84	15.8	75.2	31718	4	AAK87443	Aak87443 Human imm
C 85	15.8	75.2	31718	4	AAK87592	Aak87592 Human imm
C 86	15.8	75.2	31718	4	AAL06415	Aal06415 Human rep
C 87	15.8	75.2	31718	4	AAK06416	Aak06416 Human rep
C 88	15.8	75.2	31718	5	AAS39916	Aas39916 Genomic s
C 89	15.8	75.2	31718	5	AAS39915	Aas39915 Genomic s
C 90	15.8	75.2	31718	9	ADB32875	Adi32875 Human nov
C 91	15.8	75.2	31718	9	ADB32876	Adi32876 Human nov
C 92	15.8	75.2	31718	12	ADN41665	Adn41665 Novel hum
C 93	15.8	75.2	31718	12	ADN41666	Adn41666 Novel hum

C 94 15.8 75.2 63609 12 ADQ97537
 95 15.8 75.2 110000 9 ADBI2064 07
 96 15.8 75.2 122923 11 ACN44026
 C 97 15.8 75.2 122923 11 ACN44026
 98 15.4 73.3 170170 10 ADL13643
 99 15.4 73.3 20 AAD34672
 C 99 15.4 73.3 327 6 AAD34632
 100 15.4 73.3 640 12 ADJ75740

Adq97537 Human can
 Continuation (8 of
 Acn44026 Human gen
 Adl13643 Osteoarthritis
 Aad34672 DST CHS1
 Aad34632 HBV infec
 Adj75740 Marker ge

ALIGNMENTS

RESULT 1
 ACC80536/c

ID ACC80536 standard; DNA; 21 BP.

XX
 AC ACC80536;

DT 29-AUG-2003 (first entry)

XX Exemplary sequence for method detecting HBV DNA in a sample.

DE Hepatitis B virus; diagnosis; nucleic acid assay; ss;
 KW transcription-mediated amplification.

XX Hepatitis B virus.

XX WO2003031934-A2.

XX 17-APR-2003.

XX 09-OCT-2002; 2002WO-US032367.

XX 09-OCT-2001; 2001US-0328492P.

PR 29-MAR-2002; 2002US-0368823P.

PR 02-JUL-2002; 2002US-0393561P.

XX (CHIR) CHIRON CORP.

PA Shyamala V;

PI WPI; 2003-403124/38.

XX New isolated hepatitis B virus (HBV) capture oligonucleotides, useful for

PT detecting HBV infection in a biological sample, or in capturing HBV

PT nucleic acids.

XX Disclosure; Page 22; 50pp; English.

XX The invention relates to a novel method of detecting hepatitis B virus

CC (HBV) infections in e.g. blood samples from donors, by capturing and

CC amplifying conserved regions of the HBV genome using a transcription-

CC mediated amplification (TMA) method as well as a 5' nuclease assay. This

CC sequence represents an exemplary replacement sequence for a target

CC sequence (ACC80535) used in an internal control for the method of the

CC invention. The new method is very sensitive and is able to detect about

CC 100 infectious units (IU) of HBV in a viremic sample

XX
 SQ Sequence 21 BP; 5 A; 6 C; 5 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 8; Length 21;

Best Local Similarity 100.0%; Pred. No. 2.5;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTAGCT 21

Db 21 CAGTGACATGCAGGCTAGCT 1

RESULT 2

ACC80538

ID ACC80538 standard; DNA; 21 BP.

XX

AC ACC80538;

XX 29-AUG-2003 (first entry)

DT Internal control primer #2 for hepatitis B virus DNA detection method.

DE Primer; PCR; ss; hepatitis B virus; diagnosis; nucleic acid assay;
 KW transcription-mediated amplification.

XX Hepatitis B virus.

XX Key misc_difference 1 Location/Qualifiers

FT /tag= a

FT /note= "linked to 6-FAM"

FT misc_difference 21

FT /tag= b

FT /note= "linked to TAMRA"

XX WO2003031934-A2.

XX 17-APR-2003.

XX 09-OCT-2002; 2002WO-US032367.

XX 09-OCT-2001; 2001US-0328492P.

PR 29-MAR-2002; 2002US-0368823P.

PR 02-JUL-2002; 2002US-0393561P.

XX (CHIR) CHIRON CORP.

XX Shyamala V;

XX WPI; 2003-403124/38.

XX New isolated hepatitis B virus (HBV) capture oligonucleotides, useful for

PT detecting HBV infection in a biological sample, or in capturing HBV

PT nucleic acids.

XX Claim 14; Fig 2; 50pp; English.

XX The invention relates to a novel method of detecting hepatitis B virus

CC (HBV) infections in e.g. blood samples from donors, by capturing and

CC amplifying conserved regions of the HBV genome using a transcription-

CC mediated amplification (TMA) method as well as a 5' nuclease assay. The

CC method may also use an internal control sequence such as ACC80539, to

CC determine the level of amplification and detection by the primers and

CC probes used in the method of the invention. This sequence represents a

CC primer used to amplify the internal control region DNA sequence. The new

CC method is very sensitive and is able to detect about 100 infectious units

CC (IU) of HBV in a viremic sample

XX
 SQ Sequence 21 BP; 5 A; 5 C; 5 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 8; Length 21;

Best Local Similarity 100.0%; Pred. No. 2.5;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTAGCT 21

Db 1 CAGTGACATGCAGGCTAGCT 21

RESULT 3

ACC80537/c

ID ACC80537 standard; DNA; 21 BP.

XX

AC ACC80537;

XX

DT 29-AUG-2003 (first entry)

XX Internal control primer #1 for hepatitis B virus DNA detection method.

XX

KW Primer; PCR; ss; hepatitis B virus; diagnosis; nuclease assay;
 KW transcription-mediated amplification.

XX Hepatitis B virus.

XX Key Location/Qualifiers

FT misc_difference 1 /*tag= a

FT /note= "linked to 6-FAM"

FT misc_difference 21

FT /*tag= b

FT /note= "linked to TAMRA"

XX WO2003031934-A2.

XX 17-APR-2003.

XX 09-OCT-2002; 2002WO-US032367.

XX 09-OCT-2001; 2001US-0328492P.

XX 29-MAR-2002; 2002US-0368823P.

XX 02-JUL-2002; 2002US-0393561P.

XX (CHIR) CHIRON CORP.

XX Shyamala V;

XX WPI; 2003-403124/38.

XX New isolated hepatitis B virus (HBV) capture oligonucleotides, useful for
 PT detecting HBV infection in a biological sample, or in capturing HBV
 PT nucleic acids.

XX Claim 13; Fig 2; 50pp; English.

XX The invention relates to a novel method of detecting hepatitis B virus
 CC (HBV) infections in e.g. blood samples from donors, by capturing and
 CC amplifying conserved regions of the HBV genome using a transcription-
 CC mediated amplification (TMA) method as well as a 5' nuclease assay. The
 CC method may also use an internal control sequence such as ACC80539, to
 CC determine the level of amplification and detection by the primers and
 CC probes used in the method of the invention. This sequence represents a
 CC primer used to amplify the internal control region DNA sequence. The new
 CC method is very sensitive and is able to detect about 100 infectious units
 CC (IU) of HBV in a viremic sample

XX Sequence 21 BP; 5 A; 6 C; 5 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 8; Length 21;

Best Local Similarity 100.0%; Pred. No. 2.5;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCTAGCT 21

DB 21 CAGTGACATGCAGGTCTAGCT 1

RESULT 4

ABZ59632/c

ID ABZ59632 standard; DNA; 21 BP.

XX AC ABZ59632;

XX 22-APR-2003 (first entry)

XX Parvovirus B19 related internal control oligonucleotide SEQ ID NO:90.
 DE Human parvovirus B19; parvovirus B19; infection; virus; blood; plasma;
 KW PCR primer; ss.

XX B19 virus.
 OS Synthetic.

XX Key

XX Location/Qualifiers

XX modified_base 1

PN WO2003002753-A2.

XX 09-JAN-2003.

XX 28-JUN-2002; 2002WO-US020684.

XX 28-JUN-2001; 2001US-0302077P.

XX 19-MAR-2002; 2002US-0365956P.

XX 29-MAR-2002; 2002US-0369224P.

XX (CHIR) CHIRON CORP.

XX Pichuantes S, Shyamala V;

XX WPI; 2003-201510/19.

XX Detecting a human parvovirus B19 infection in a biological sample to
 PT prevent viral transmission, comprises reacting a parvovirus B19 nucleic
 PT acid with a primer complementary to the 3'-terminal portion of the RNA
 PT target sequence.

XX Claim 8; Page 29; 148pp; English.

XX The present invention describes a method for detecting a human parvovirus
 CC B19 infection in a biological sample. The method comprises reacting the
 CC isolated parvovirus B19 nucleic acid with a first oligonucleotide
 CC consisting of a first primer containing a complexing sequence
 CC sufficiently complementary to the 3'-terminal portion of the RNA target
 CC sequence to complex with. Also described: (1) amplifying a target
 CC parvovirus B19 nucleotide sequence; (2) a polynucleotide comprising one
 CC of 47 700 base pair sequences (see ABZ59549 to ABZ59569, and ABZ59604 to
 CC ABZ59629); (3) a polynucleotide comprising either of 2 4678 base pair
 CC sequences (see ABZ59570 and ABZ59571); (4) an oligonucleotide primer
 CC consisting of a promoter region recognised by a DNA-dependent RNA
 CC polymerase operably linked to a human parvovirus B19-specific complexing
 CC sequence of 10-75 nucleotides; (5) an oligonucleotide probe comprising a
 CC parvovirus B19-specific hybridising sequence of 10-50 nucleotides linked
 CC to an acridinium ester label; and (6) a diagnostic test kit comprising an
 CC oligonucleotide primer of (4), and instructions for conducting the
 CC diagnostic test. The method is useful for detecting parvovirus infection
 CC in a biological sample, such as in blood products, to prevent
 CC transmission of the virus through blood and plasma derivatives or by
 CC close personal contact. ABZ59549 to ABZ59634 and ABP57262 to ABP57267
 CC represent sequences used in the exemplification of the present invention

XX Sequence 21 BP; 5 A; 6 C; 5 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 9; Length 21;

Best Local Similarity 100.0%; Pred. No. 2.5;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCTAGCT 21

DB 21 CAGTGACATGCAGGTCTAGCT 1

RESULT 5

ADI53796

ID ADI53796 standard; DNA; 21 BP.

XX AC ADI53796;

XX 06-MAY-2004 (first entry)

XX HAV internal control specific detection probe.
 DE HAV; nucleic acid amplification; nucleic acid detection; ss; probe.

XX Hepatitis A virus.

OS Synthetic.

XX Key

XX Location/Qualifiers

XX modified_base 1

```

FT FT /*tag= a
FT FT /mod_base= 5'-TET labelled
FT FT 21
FT FT /*tag= b
FT FT /mod_base= 3'-TAMRA labelled
XX
XX WO2003106641-A2.
XX
XX 24-DEC-2003.
XX
XX 12-JUN-2003; 2003WO-US018827.
XX
XX 12-JUN-2002; 2002US-0388544P.
XX
XX (CHIR ) CHIRON CORP.
XX
XX Shyamala V;
XX
XX WPI; 2004-082181/08.
XX
XX Hepatitis A virus specific primers and probes derived from conserved
XX PT regions of the hepatitis A virus genome, useful in nucleic acid-based
XX PT diagnostic tests for the detection of Hepatitis A virus in biological
XX PT samples.
XX
XX PA (CHIR ) CHIRON CORP.
XX
XX PI Shyamala V;
XX
XX DR WPI; 2004-082181/08.
XX
XX PT Hepatitis A virus specific primers and probes derived from conserved
XX PT regions of the hepatitis A virus genome, useful in nucleic acid-based
XX PT diagnostic tests for the detection of Hepatitis A virus in biological
XX PT samples.
XX
XX PS Example 2; SEQ ID NO 18; 44pp; English.
XX
XX CC The invention relates to Hepatitis A virus (HAV) specific primers and
XX CC probes derived from conserved regions of the hepatitis A virus genome.
XX CC The HAV-specific primers and probes are used in a method for detecting
XX CC HAV in a biological sample. Also provided are capture oligonucleotides
XX CC {Seq ID. No. 10}-{Seq ID. No. 14} which are used in a method for
XX CC detecting HAV infection in a biological sample. The present sequence
XX CC represents a detection probe specific for an internal control sequence.
XX
XX SQ Sequence 21 BP; 5 A; 5 C; 6 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 12; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.5;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTCTAGCT 21
Db 1 CAGTGACATGCAGGCTCTAGCT 21

RESULT 6
AD153797
ID AD153797 standard; DNA; 23 BP.
XX
XX AC AD153797;
XX
XX 06-MAY-2004 (first entry)
XX
XX HAV internal control specific detection probe.
XX
XX HAV; nucleic acid amplification; nucleic acid detection; ss; probe.
XX
XX Hepatitis A virus.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
FH modified_base 1 /*tag= a
FT FT /mod_base= 5'-TET labelled
FT FT 23
FT FT /*tag= b
FT FT /mod_base= 3'-TAMRA labelled
XX
XX WO2003106641-A2.
XX
XX 24-DEC-2003.
XX

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PF 12-JUN-2003; 2003WO-US018827.
XX
PR 12-JUN-2002; 2002US-0388544P.
XX
XX (CHIR ) CHIRON CORP.
XX
XX Shyamala V;
XX
XX WPI; 2004-082181/08.
XX
XX Hepatitis A virus specific primers and probes derived from conserved
XX PT regions of the hepatitis A virus genome, useful in nucleic acid-based
XX PT diagnostic tests for the detection of Hepatitis A virus in biological
XX PT samples.
XX
XX PS Example 2; SEQ ID NO 19; 44pp; English.
XX
XX CC The invention relates to Hepatitis A virus (HAV) specific primers and
XX CC probes derived from conserved regions of the hepatitis A virus genome.
XX CC The HAV-specific primers and probes are used in a method for detecting
XX CC HAV in a biological sample. Also provided are capture oligonucleotides
XX CC {Seq ID. No. 10}-{Seq ID. No. 14} which are used in a method for
XX CC detecting HAV infection in a biological sample. The present sequence
XX CC represents a detection probe specific for an internal control sequence.
XX
XX SQ Sequence 23 BP; 5 A; 7 C; 6 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 12; Length 23;
Best Local Similarity 100.0%; Pred. No. 2.5;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTCTAGCT 21
Db 3 CAGTGACATGCAGGCTCTAGCT 23

RESULT 7
ADQ30670
ID ADQ30670 standard; DNA; 23 BP.
XX
XX AC ADQ30670;
XX
XX 23-SEP-2004 (first entry)
XX
XX West Nile Virus internal control probe #1.
XX
XX ss; primer; West Nile Virus; diagnosis.
XX
XX West Nile virus.
XX
XX WO2004055159-A2.
XX
XX 01-JUL-2004.
XX
XX 05-DEC-2003; 2003WO-US038750.
XX
XX 12-DEC-2002; 2002US-0432850P.
XX
XX 20-JUN-2003; 2003US-0480431P.
XX
XX (CHIR ) CHIRON CORP.
XX
XX Shyamala V;
XX
XX WPI; 2004-488058/46.
XX
XX New isolated oligonucleotides for accurately diagnosing West Nile virus
XX PT infection or for capturing, detecting and quantitating West Nile virus in
XX PT blood samples.
XX
XX Claim 29; SEQ ID NO 40; 56pp; English.
XX
XX The invention relates to an isolated oligonucleotide not more than 60
XX CC nucleotides in length comprising a nucleotide sequence (S1) of at least
XX CC

```


CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
CC 20, 21 or 23 bp) given in the specification derived from the West Nile
CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
CC identity to the nucleotide sequence of (S1), or complements of (S1) and
CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
CC end and/or the 3'-end. The detectable label is a fluorescent label
CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
CC composition and methods are useful for accurately diagnosing West Nile
CC virus infection or for capturing, detecting and quantitating West Nile
CC virus in biological samples, particularly blood samples. This sequence
CC corresponds to a probe to the internal control sequence for the detection
CC of WNV sequences using the oligonucleotides of the invention.

XX
SQ Sequence 23 BP; 5 A; 5 C; 6 G; 5 T; 0 U; 2 Other;

Query Match 100.0%; Score 21; DB 12; Length 23;
Best Local Similarity 100.0%; Pred. No. 2.5;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGCTCTAGCT 21
|||||
Db 2 CAGTGACATGCAGGCTCTAGCT 22

RESULT 8

ADQ30671

ID ADQ30671 standard; DNA; 25 BP. -

AC ADQ30671;

XX

XX 23-SEP-2004 (first entry)

DT

DE West Nile Virus internal control probe #2.

XX

XX ss; primer; West Nile Virus; diagnosis.

KW

XX West Nile virus.

OS

XX WO2004055159-A2.

PN

XX 01-JUL-2004.

PD

XX 05-DEC-2003; 2003WO-US038750.

PF

XX 12-DEC-2002; 2002US-0432850P.

PR

PR 20-JUN-2003; 2003US-0480431P.

XX

XX (CHIR) CHIRON CORP.

PA

XX Shyamala V;

PI

XX WPI; 2004-488058/46.

DR

XX New isolated oligonucleotides for accurately diagnosing West Nile virus

PT infection or for capturing, detecting and quantitating West Nile virus in

PT blood samples.

XX

PS Claim 29; SEQ ID NO 41; 56pp; English.

XX

CC The invention relates to an isolated oligonucleotide not more than 60
CC nucleotides in length comprising a nucleotide sequence (S1) of at least
CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
CC 20, 21 or 23 bp) given in the specification derived from the West Nile
CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
CC identity to the nucleotide sequence of (S1), or complements of (S1) and
CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
CC end and/or the 3'-end. The detectable label is a fluorescent label
CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
CC composition and methods are useful for accurately diagnosing West Nile
CC virus infection or for capturing, detecting and quantitating West Nile
CC virus in biological samples, particularly blood samples. This sequence

CC corresponds to a probe to the internal control sequence for the detection
CC of WNV sequences using the oligonucleotides of the invention.

XX
SQ Sequence 25 BP; 5 A; 7 C; 6 G; 5 T; 0 U; 2 Other;

Query Match 100.0%; Score 21; DB 12; Length 25;
Best Local Similarity 100.0%; Pred. No. 2.5;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGCTCTAGCT 21
|||||
Db 4 CAGTGACATGCAGGCTCTAGCT 24

RESULT 9

ABZ59634

ID ABZ59634 standard; DNA; 681 BP.

XX

XX AC ABZ59634;

XX

XX 22-APR-2003 (first entry)

DT

DE Exemplary internal control nucleotide sequence SEQ ID NO:92.

XX

XX Human parvovirus B19; parvovirus B19; infection; virus; blood; plasma;

KW

XX gene; ds.

XX

XX Synthetic.

OS

XX WO2003002753-A2.

PN

XX 09-JAN-2003.

PD

XX 28-JUN-2002; 2002WO-US020684.

PF

XX 28-JUN-2001; 2001US-0302077P.

PR

PR 19-MAR-2002; 2002US-0365956P.

XX

XX 29-MAR-2002; 2002US-0369224P.

XX

XX (CHIR) CHIRON CORP.

PA

XX Pichuanes S, Shyamala V;

PI

XX WPI; 2003-201510/19.

DR

XX Detecting a human parvovirus B19 infection in a biological sample to

PT prevent viral transmission, comprises reacting a parvovirus B19 nucleic

PT acid with a primer complementary to the 3'-terminal portion of the RNA

PT target sequence.

XX

XX Claim 7; Fig 12; 148pp; English.

PS

XX The present invention describes a method for detecting a human parvovirus

CC B19 infection in a biological sample. The method comprises reacting the

CC isolated parvovirus B19 nucleic acid with a first oligonucleotide

CC consisting of a first primer containing a complexing sequence

CC sufficiently complementary to the 3'-terminal portion of the RNA target

CC sequence to complex with. Also described: (1) amplifying a target

CC parvovirus B19 nucleotide sequence; (2) a polynucleotide comprising one

CC of 47 700 base pair sequences (see ABZ59549 to ABZ59569, and ABZ59604 to

CC ABZ59629); (3) a polynucleotide comprising either of 2 4678 base pair

CC sequences (see ABZ59570 and ABZ59571); (4) an oligonucleotide primer

CC consisting of a promoter region recognised by a DNA-dependent RNA

CC polymerase operably linked to a human parvovirus B19-specific complexing

CC sequence of 10-75 nucleotides; (5) an oligonucleotide probe comprising a

CC parvovirus B19-specific hybridising sequence of 10-50 nucleotides linked

CC to an acridinium ester label; and (6) a diagnostic test kit comprising an

CC oligonucleotide primer of (4), and instructions for conducting the

CC diagnostic test. The method is useful for detecting parvovirus infection

CC in a biological sample, such as in blood products, to prevent

CC transmission of the virus through blood and plasma derivatives or by

CC close personal contact. ABZ59549 to ABZ59634 and ABP57262 to ABP57267

CC represent sequences used in the exemplification of the present invention

```
XX SQ Sequence 681 BP; 206 A; 138 C; 137 G; 200 T; 0 U; 0 Other;
Query Match 100.0%; Score 21; DB 9; Length 681;
Best Local Similarity 100.0%; Pred. No. 3.4;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTTAGCT 21
Db 62 CAGTGACATGCAGGTTAGCT 82

RESULT 10
ADIS3795
ID ADIS3795 standard; DNA; 727 BP.
XX AC
ADIS3795;
XX DT 06-MAY-2004 (first entry)
XX DE HAV internal control sequence.
XX KW HAV; nucleic acid amplification; nucleic acid detection; ds.
XX OS Hepatitis A virus.
XX OS Synthetic.
XX PN WO2003106641-A2.
XX PD 24-DEC-2003.
XX PF 12-JUN-2003; 2003WO-US018827.
XX PR 12-JUN-2002; 2002US-0388544P.
XX PA (CHIR ) CHIRON CORP.
XX PI Shyamala V;
XX DR WPI; 2004-082181/08.
XX PT Hepatitis A virus specific primers and probes derived from conserved
PT regions of the hepatitis A virus genome, useful in nucleic acid-based
PT diagnostic tests for the detection of Hepatitis A virus in biological
PT samples.
XX PS Example 2; SEQ ID NO 17; 44pp; English.
XX CC The invention relates to Hepatitis A virus (HAV) specific primers and
CC probes derived from conserved regions of the hepatitis A virus genome.
CC The HAV-specific primers and probes are used in a method for detecting
CC HAV in a biological sample. Also provided are capture oligonucleotides
CC {Seq ID. No. 10}-{Seq ID. No. 14} which are used in a method for
CC detecting HAV infection in a biological sample. The present sequence
CC represents an internal control sequence used as a control for target
CC capture and amplification.
XX SQ Sequence 727 BP; 147 A; 169 C; 186 G; 225 T; 0 U; 0 Other;
Query Match 100.0%; Score 21; DB 12; Length 727;
Best Local Similarity 100.0%; Pred. No. 3.4;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTTAGCT 21
Db 581 CAGTGACATGCAGGTTAGCT 601

RESULT 11
ADQ30647
ID ADQ30647 standard; DNA; 967 BP.
XX AC
ADQ30647;
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XX DT 23-SEP-2004 (first entry)
XX DE West Nile virus internal diagnosis control sequence.
XX KW ss; internal control; West Nile Virus; diagnosis.
XX OS West Nile virus.
XX PN WO2004055159-A2.
XX PD 01-JUL-2004.
XX PF 05-DEC-2003; 2003WO-US038750.
XX PR 12-DEC-2002; 2002US-0432850P.
XX PR 20-JUN-2003; 2003US-0480431P.
XX PA (CHIR ) CHIRON CORP.
XX PI Shyamala V;
XX DR WPI; 2004-488058/46.
XX PT New isolated oligonucleotides for accurately diagnosing West Nile virus
PT infection or for capturing, detecting and quantitating West Nile virus in
PT blood samples.
XX PS Claim 27; SEQ ID NO 17; 56pp; English.
XX CC The invention relates to an isolated oligonucleotide not more than 60
CC nucleotides in length comprising a nucleotide sequence (S1) of at least
CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
CC 20, 21 or 23 bp) given in the specification derived from the West Nile
CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
CC identity to the nucleotide sequence of (S1), or complements of (S1) and
CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
CC end and/or the 3'-end. The detectable label is a fluorescent label
CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
CC composition and methods are useful for accurately diagnosing West Nile
CC virus infection or for capturing, detecting and quantitating West Nile
CC virus in biological samples, particularly blood samples. This sequence
CC corresponds to an internal control sequence for the detection of WNV
CC sequences using the oligonucleotides of the invention.
XX SQ Sequence 967 BP; 273 A; 206 C; 272 G; 216 T; 0 U; 0 Other;
Query Match 100.0%; Score 21; DB 12; Length 967;
Best Local Similarity 100.0%; Pred. No. 3.5;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTTAGCT 21
Db 153 CAGTGACATGCAGGTTAGCT 173

RESULT 12
ACC80539
ID ACC80539 standard; DNA; 1696 BP.
XX AC
ACC80539;
XX DT 29-AUG-2003 (first entry)
XX DE Internal control region for hepatitis B virus DNA detection method.
XX KW Hepatitis B virus; diagnosis; nucleic acid assay; internal control region;
KW transcription-mediated amplification; ds.
XX OS Hepatitis B virus.
XX PN WO2003031934-A2.
```

XX PD 17-APR-2003.
 XX PF 09-OCT-2002; 2002WO-US032367.
 XX PR 09-OCT-2001; 2001US-0328492P.
 PR 29-MAR-2002; 2002US-0368823P.
 PR 02-JUL-2002; 2002US-0393561P.
 XX PA (CHIR) CHIRON CORP.
 XX PI Shyamala V;
 XX PF WI; 2003-403124/38.
 XX PT New isolated hepatitis B virus (HBV) capture oligonucleotides, useful for
 PT detecting HBV infection in a biological sample, or in capturing HBV
 PT nucleic acids.
 XX PS Disclosure; Fig 3; 50pp; English.
 XX CC The invention relates to a novel method of detecting hepatitis B virus
 CC (HBV) infections in e.g. blood samples from donors, by capturing and
 CC amplifying conserved regions of the HBV genome using a transcription-
 CC mediated amplification (TMA) method as well as a 5' nuclease assay. The
 CC method may also use an internal control sequence (this sequence), to
 CC determine the level of amplification and detection by the primers and
 CC probes used in the method of the invention. The new method is very
 CC sensitive and is able to detect about 100 infectious units (IU) of HBV in
 CC a viremic sample
 XX SQ Sequence 1696 BP; 359 A; 462 C; 386 G; 489 T; 0 U; 0 Other;
 Query Match 100.0%; Score 21; DB 8; Length 1696;
 Best Local Similarity 100.0%; Pred. No. 3.7;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 CAGTGACATGCAGGCTCTAGCT 21
 Db 1428 CAGTGACATGCAGGCTCTAGCT 1448
 RESULT 13
 ADE711204/c
 ID ADE71204 standard; DNA; 5446 BP.
 AC ADE71204;
 XX 29-JAN-2004 (first entry)
 DT Novel human protein coding sequence #20.
 DE human; novel protein; drug; gene; ds.
 KW Homo sapiens.
 XX JP2002345493-A.
 XX 03-DEC-2002.
 PD 29-MAR-2001; 2002JP-00049046.
 PF 29-MAR-2001; 2001JP-00095524.
 XX (KAZU-) ZH KAZUSA DNA KENKYUSHO.
 XX WI; 2003-460885/44.
 DR P-PSDB; ADE71266.
 XX A gene and a protein encoded by it, used in drugs.
 PT Claim 1; SEQ ID NO 20; 257pp; Japanese.
 XX

CC The invention comprises the amino acid and coding sequences of novel
 CC human proteins. The DNA and protein sequences of the invention are used
 CC in drugs. The present DNA sequence encodes a novel human protein of the
 CC invention.
 XX SQ Sequence 5446 BP; 1477 A; 1320 C; 1292 G; 1357 T; 0 U; 0 Other;
 Query Match 82.9%; Score 17.4; DB 10; Length 5446;
 Best Local Similarity 94.7%; Pred. No. 2.3e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 OY 1 CAGTGACATGCAGGCTCTAG 19
 Db 833 CAGTGACATGCAGGCTCTAG 815
 RESULT 14
 ABZ73855/c
 ID ABZ73855 standard; DNA; 28482 BP.
 XX AC ABZ73855;
 XX 12-MAY-2003 (first entry)
 DT Secreted protein gene 64 genomic fragment HCEGX05, SEQ ID NO:1002.
 DE Human; secreted protein; cancer; tumour; hyperproliferative disorder;
 KW autoimmune disorder; inflammation; angiogenic diseases; AIDS;
 KW acquired immunodeficiency syndrome; hepatitis; anaemia; wound healing;
 KW drug screening; chromosome identification; chromosome mapping;
 KW cytotoxic; gene therapy; antiinflammatory; immunomodulator; anti-HIV;
 KW antianaemic; vulnery; chromosome 20q13; gene; ds.
 XX OS Homo sapiens.
 XX WO200277013-A2.
 XX 03-OCT-2002.
 XX 26-MAR-2002; 2002WO-US009370.
 XX 27-MAR-2001; 2001US-0278650P.
 PR 12-SEP-2001; 2001US-00950082.
 PR 12-SEP-2001; 2001US-00950083.
 XX (HUMA-) HUMAN GENOME SCI INC.
 XX PI Rosen CA, Ruben SM;
 XX WI; 2003-040578/03.
 XX New human secreted proteins and nucleic acids, useful for detecting or
 PT treating cancer or other hyperproliferative disorders, autoimmune
 PT disorders, inflammatory disorders, HIV disease, hepatitis or anemia.
 XX Disclosure; Page 1651-1658; 2474pp; English.
 XX ABZ73281-ABZ73697 represent cDNAs corresponding to 391 human secreted
 CC protein genes, and ABP00547-ABP01363 represent the proteins they encode.
 CC ABZ73698-ABZ74687 represent human secreted protein genomic fragments. The
 CC invention also encompasses antibodies specific for the secreted proteins,
 CC the use of the secreted proteins in drug screening and recombinant
 CC vectors and host cells comprising a nucleic acid of the invention. The
 CC secreted proteins are thought to be involved in biological activities
 CC associated with cellular signalling, cellular differentiation, cell
 CC migration, prohormone activation and neurotransmitter activity. The
 CC secreted proteins, nucleic acids encoding them, antibodies or antibody
 CC fragments specific for the secreted proteins, and modulators of protein
 CC activity are useful for diagnosing or treating cancers or other
 CC hyperproliferative disorders. Additionally, the secreted proteins and
 CC their nucleic acids may also be used in the treatment of autoimmune
 CC disorders, inflammatory disorders, diseases involving angiogenesis, AIDS
 CC (acquired immunodeficiency syndrome), hepatitis, anaemia, and to promote

CC wound healing. Nucleic acids of the invention may be used for chromosome
 CC identification, chromosome mapping, in gene therapy, for identifying
 CC individuals from minute biological samples, as hybridisation probes, and
 CC as molecular weight markers. The present sequence represents a human
 CC secreted protein genomic fragment referred to in the disclosure of the
 CC invention
 XX
 SQ Sequence 28482 BP; 7636 A; 6245 C; 6763 G; 7838 T; 0 U; 0 Other;
 Query Match 82.9%; Score 17.4; DB 8; Length 28482;
 Best Local Similarity 94.7%; Pred. No. 2.6e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 CAGTGACATGCAGGTCTAG 19
 ||||| ||||| ||||| ||||| |||||
 Db 12331 CAGTGACATGCAGGTCTAG 12313
 RESULT 15
 ADA44262/c
 ID ADA44262 standard; DNA; 28482 BP.
 AC ADA44262;
 XX
 DT 20-NOV-2003 (first entry)
 XX
 DE Human secreted protein DNA SEQ ID 455.
 XX
 KW Gene therapy; human; Antidiabetic; Anorectic; Ophthalmological;
 KW Neuroprotective; Cerebroprotective; Antianemic; ds.
 XX
 OS Homo sapiens.
 XX
 PN WO200300865-A2.
 XX
 PD 03-JAN-2003.
 XX
 PF 26-MAR-2002; 2002WO-US009105.
 XX
 PR 27-MAR-2001; 2001US-0278650P.
 PR 12-SEP-2001; 2001US-00950082.
 PR 12-SEP-2001; 2001US-00950083.
 XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 PI Rosen CA, Ruben SM;
 XX
 DR WPI; 2003-184045/18.
 XX
 PT A human secreted protein and nucleic acids useful for preparing a
 PT diagnostic or pharmaceutical composition for diagnosing or treating
 PT diabetes or conditions related to diabetes, e.g. hyperglycemia, obesity,
 PT retinopathy, neuropathy.
 XX
 PS Disclosure; SEQ ID NO 455; 701pp; English.
 XX
 CC The invention relates to novel genes and their fragments which are useful
 CC for preventing, treating or ameliorating medical conditions e.g. by
 CC protein or gene therapy. The genes are isolated from a range of human
 CC tissues disclosed in the specification. The nucleic acids and proteins
 CC are useful in the diagnosis, treatment and prevention of conditions
 CC related to diabetes, e.g. hyperglycaemia, obesity, retinopathy,
 CC polynuropathy, atherosclerosis, anaemia, stroke, gangrene, impotence,
 CC infection, cataract, renal disorders, or endocrine disorders. The present
 CC sequence was used to illustrate the invention.
 XX
 SQ Sequence 28482 BP; 7636 A; 6245 C; 6763 G; 7838 T; 0 U; 0 Other;
 Query Match 82.9%; Score 17.4; DB 8; Length 28482;
 Best Local Similarity 94.7%; Pred. No. 2.6e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 CAGTGACATGCAGGTCTAG 19

Db 12331 CAGTGACATGCAGGTCTAG 12313
 ||||| ||||| ||||| ||||| |||||
 RESULT 16
 ABZ74517/c
 ID ABZ74517 standard; DNA; 32681 BP.
 XX
 AC ABZ74517;
 XX
 DT 12-MAY-2003 (first entry)
 XX
 DE Secreted protein gene 329 genomic fragment HTLBT80, SEQ ID NO:1664.
 XX
 KW Human; secreted protein; cancer; tumour; hyperproliferative disorder;
 KW autoimmune disorder; inflammation; angiogenic diseases; AIDS;
 KW acquired immunodeficiency syndrome; hepatitis; anaemia; wound healing;
 KW drug screening; chromosome identification; chromosome mapping;
 KW cytostatic; gene therapy; antiinflammatory; immunomodulator; anti-HIV;
 KW antianaemic; vulnery; chromosome 20q11.21-13.11; gene; ds.
 XX
 OS Homo sapiens.
 XX
 PN WO200277013-A2.
 XX
 PD 03-OCT-2002.
 XX
 PF 26-MAR-2002; 2002WO-US009370.
 XX
 PR 27-MAR-2001; 2001US-0278650P.
 PR 12-SEP-2001; 2001US-00950082.
 PR 12-SEP-2001; 2001US-00950083.
 XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 PI Rosen CA, Ruben SM;
 XX
 DR WPI; 2003-040578/03.
 XX
 PT New human secreted proteins and nucleic acids, useful for detecting or
 PT treating cancer or other hyperproliferative disorders, autoimmune
 PT disorders, inflammatory disorders, HIV disease, hepatitis or anemia.
 XX
 PS Disclosure; Page 2245-2253; 247app; English.
 XX
 CC ABZ73281-ABZ73697 represent cDNAs corresponding to 391 human secreted
 CC protein genes, and ABP00947-ABP01363 represent the proteins they encode.
 CC ABZ73698-ABZ74687 represent human secreted protein genomic fragments. The
 CC invention also encompasses antibodies specific for the secreted proteins,
 CC the use of the secreted proteins in drug screening and recombinant
 CC vectors and host cells comprising a nucleic acid of the invention. The
 CC secreted proteins are thought to be involved in biological activities
 CC associated with cellular signalling, cellular differentiation, cell
 CC migration, prohormone activation and neurotransmitter activity. The
 CC secreted proteins, nucleic acids encoding them, antibodies or antibody
 CC fragments specific for the secreted proteins, and modulators of protein
 CC activity are useful for diagnosing or treating cancers or other
 CC hyperproliferative disorders. Additionally, the secreted proteins and
 CC their nucleic acids may also be used in the treatment of autoimmune
 CC disorders, inflammatory disorders, diseases involving angiogenesis, AIDS
 CC (acquired immunodeficiency syndrome), hepatitis, anaemia, and to promote
 CC wound healing. Nucleic acids of the invention may be used for chromosome
 CC identification, chromosome mapping, in gene therapy, for identifying
 CC individuals from minute biological samples, as hybridisation probes, and
 CC as molecular weight markers. The present sequence represents a human
 CC secreted protein genomic fragment referred to in the disclosure of the
 CC invention
 XX
 SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;
 Query Match 82.9%; Score 17.4; DB 8; Length 32681;
 Best Local Similarity 94.7%; Pred. No. 2.7e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCTAG 19
|||||
Db 16526 CAGTGACAGCGAGGTCTAG 16508

RESULT 17
ABZ73854/c
ID ABZ73854 standard; DNA; 32681 BP.
XX
AC ABZ73854;
XX
DT 12-MAY-2003 (first entry)
XX
DE Secreted protein gene 64 genomic fragment HCEGX05, SEQ ID NO:1001.
XX
KW Human; secreted protein; cancer; tumour; hyperproliferative disorder;
KW autoimmune disorder; inflammation; angioinflammatory diseases; AIDS;
KW acquired immunodeficiency syndrome; hepatitis; anaemia; wound healing;
KW drug screening; chromosome identification; chromosome mapping;
KW cytostatic; gene therapy; antiinflammatory; immunomodulator; anti-HIV;
KW antianaemic; vulnery; chromosome 20q13; gene; ds.
XX
OS Homo sapiens.
XX
PN WO200277013-A2.
XX
PD 03-OCT-2002.
XX
PF 26-MAR-2002; 2002WO-US009370.
XX
PR 27-MAR-2001; 2001US-0278650P.
PR 12-SEP-2001; 2001US-00950082.
PR 12-SEP-2001; 2001US-00950083.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Ruben SM;
PI WPI; 2003-040578/03.
XX
XX New human secreted proteins and nucleic acids, useful for detecting or
PT treating cancer or other hyperproliferative disorders, autoimmune
PT disorders, inflammatory disorders, HIV disease, hepatitis or anemia.
XX
PS Disclosure; Page 1643-1651; 2474pp; English.
XX
XX ABZ73281-ABZ73697 represent cDNAs corresponding to 391 human secreted
CC protein genes, and ABP00947-ABP01363 represent the proteins they encode.
CC ABZ73698-ABZ74687 represent human secreted protein genomic fragments. The
CC invention also encompasses antibodies specific for the secreted proteins,
CC the use of the secreted proteins in drug screening and recombinant
CC vectors and host cells comprising a nucleic acid of the invention. The
CC secreted proteins are thought to be involved in biological activities
CC associated with cellular signalling, cellular differentiation, cell
CC migration, pro-hormone activation and neurotransmitter activity. The
CC secreted proteins, nucleic acids encoding them, antibodies or antibody
CC fragments specific for the secreted proteins, and modulators of protein
CC activity are useful for diagnosing or treating cancers or other
CC hyperproliferative disorders. Additionally, the secreted proteins and
CC their nucleic acids may also be used in the treatment of autoimmune
CC disorders, inflammatory disorders, diseases involving angiogenesis, AIDS
CC (acquired immunodeficiency syndrome), hepatitis, anaemia, and to promote
CC wound healing. Nucleic acids of the invention may be used for chromosome
CC identification, chromosome mapping, in gene therapy, for identifying
CC individuals from minute biological samples, as hybridisation probes, and
CC as molecular weight markers. The present sequence represents a human
CC secreted protein genomic fragment referred to in the disclosure of the
CC invention
XX
SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;
Query Match 82.9%; Score 17.4; DB 8; Length 32681;

Best Local Similarity 94.7%; Pred. No. 2.7e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCTAG 19
|||||
Db 16526 CAGTGACAGCGAGGTCTAG 16508

RESULT 18
ADA98915/c
ID ADA98915 standard; DNA; 32681 BP.
XX
AC ADA98915;
XX
DT 20-NOV-2003 (first entry)
XX
DE Human secreted protein-related DNA sequence #508.
XX
KW human; secreted protein; cardiovascular disorder; arrhythmia;
KW atherosclerosis; stroke; endocarditis; congestive heart failure;
KW rheumatic heart disease; cardiomyopathy; hemorrhoids; varicose veins;
KW migraine; thrombosis; neural disorder; immune system disorder;
KW muscular disorder; reproductive disorder; gastrointestinal disorder;
KW pulmonary disorder; renal disorder; proliferative disorder; cancer; ds.
XX
OS Homo sapiens.
XX
PN WO2003004623-A2.
XX
PD 16-JAN-2003.
XX
PF 26-MAR-2002; 2002WO-US009922.
XX
PR 27-MAR-2001; 2001US-0278650P.
PR 12-SEP-2001; 2001US-00950082.
PR 12-SEP-2001; 2001US-00950083.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Ruben SM;
PI WPI; 2003-247946/24.
XX
XX New human secreted polypeptide and nucleic acid molecules, useful for
PT diagnosing, preventing, prognosticating or treating cardiovascular
PT disorders (e.g. arrhythmia, atherosclerosis, cardiomyopathy, or
PT thrombosis).
XX
PS Disclosure; SEQ ID NO 1024; 1572pp; English.
XX
XX The invention comprises the amino acid and coding sequence of human
CC secreted proteins. The DNA and protein sequences of the invention are
CC useful in the treatment of cardiovascular disorders, such as: arrhythmia,
CC atherosclerosis, stroke, endocarditis, congestive heart failure,
CC rheumatic heart disease, cardiomyopathy, hemorrhoids, varicose veins,
CC migraine, or thrombosis. The DNA and protein sequences may also be used
CC for treating or preventing: neural disorders, immune system disorders,
CC muscular disorders, reproductive disorders, gastrointestinal disorders,
CC pulmonary disorders, renal disorders, proliferative disorders and/or
CC cancerous diseases. The present DNA sequence is used in the
CC exemplification of the invention. NOTE: The present sequence is shown on
CC the WIPO website.
XX
SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;
Query Match 82.9%; Score 17.4; DB 8; Length 32681;

Best Local Similarity 94.7%; Pred. No. 2.7e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCTAG 19
|||||
Db 16526 CAGTGACAGCGAGGTCTAG 16508

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RESULT 19
ADA44261/c
ID ADA44261 standard; DNA; 32681 BP.
XX AC
XX ADA44261;
XX AC
XX 20-NOV-2003 (first entry)
DT DT
XX 20-NOV-2003 (first entry)
DE DE
XX Human secreted protein DNA SEQ ID 454.
XX KW
XX Gene therapy; human; Antidiabetic; Anorectic; Ophthalmological;
XX Neuroprotective; Cerebroprotective; Antianemic; ds.
XX OS
XX Homo sapiens.
XX PN
XX WO2003000865-A2.
XX PD
XX 03-JAN-2003.
XX PF
XX 26-MAR-2002; 2002WO-US009105.
XX PR
XX 27-MAR-2001; 2001US-0278650P.
XX 12-SEP-2001; 2001US-00950082.
XX 12-SEP-2001; 2001US-00950083.
XX PA
XX (HUMA-) HUMAN GENOME SCI INC.
XX PI
XX Rosen CA, Ruben SM;
XX WPI; 2003-184045/18.
XX DR
XX 03-JAN-2003.
XX PF
XX 26-MAR-2002; 2002WO-US009105.
XX PR
XX 27-MAR-2001; 2001US-0278650P.
XX 12-SEP-2001; 2001US-00950082.
XX 12-SEP-2001; 2001US-00950083.
XX PA
XX (HUMA-) HUMAN GENOME SCI INC.
XX PI
XX Rosen CA, Ruben SM;
XX WPI; 2003-184045/18.
XX DR
XX A human secreted protein and nucleic acids useful for preparing a
PT diagnostic or pharmaceutical composition for diagnosing or treating
PT diabetes or conditions related to diabetes, e.g. hyperglycemia, obesity,
PT retinopathy, neuropathy.
XX FS
XX Disclosure; SEQ ID NO 454; 701pp; English.
XX CC
XX The invention relates to novel genes and their fragments which are useful
CC for preventing, treating or ameliorating medical conditions e.g. by
CC protein or gene therapy. The genes are isolated from a range of human
CC tissues disclosed in the specification. The nucleic acids and proteins
CC are useful in the diagnosis, treatment and prevention of conditions
CC related to diabetes, e.g. hyperglycaemia, obesity, retinopathy,
CC polynuropathy, atherosclerosis, anaemia, stroke, gangrene, impotence,
CC infection, cataract, renal disorders, or endocrine disorders. The present
CC sequence was used to illustrate the invention.
XX SQ
XX Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;

Query Match 82.9%; Score 17.4; DB 8; Length 32681;
Best Local Similarity 94.7%; Pred. No. 2.7e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTCTAG 19
Db 16526 CAGTGACAGCAGGCTCTAG 16508

RESULT 20
ADA44519/c
ID ADA44519 standard; DNA; 32681 BP.
XX AC
XX ADA44519;
XX AC
XX 20-NOV-2003 (first entry)
DT DT
XX Human secreted protein DNA SEQ ID 712.
XX DE
XX Gene therapy; human; Antidiabetic; Anorectic; Ophthalmological;
XX Neuroprotective; Cerebroprotective; Antianemic; ds.
XX KW
XX Neuroprotective; Cerebroprotective; Antianemic; ds.
XX PF
XX 26-MAR-2002; 2002WO-US009257.
XX PP

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OS Homo sapiens.
XX PN
XX WO2003000865-A2.
XX XX
XX PD
XX 03-JAN-2003.
XX PF
XX 26-MAR-2002; 2002WO-US009105.
XX PR
XX 27-MAR-2001; 2001US-0278650P.
XX 12-SEP-2001; 2001US-00950082.
XX 12-SEP-2001; 2001US-00950083.
XX PA
XX (HUMA-) HUMAN GENOME SCI INC.
XX PI
XX Rosen CA, Ruben SM;
XX WPI; 2003-184045/18.
XX DR
XX A human secreted protein and nucleic acids useful for preparing a
PT diagnostic or pharmaceutical composition for diagnosing or treating
PT diabetes or conditions related to diabetes, e.g. hyperglycemia, obesity,
PT retinopathy, neuropathy.
XX FS
XX Disclosure; SEQ ID NO 712; 701pp; English.
XX CC
XX The invention relates to novel genes and their fragments which are useful
CC for preventing, treating or ameliorating medical conditions e.g. by
CC protein or gene therapy. The genes are isolated from a range of human
CC tissues disclosed in the specification. The nucleic acids and proteins
CC are useful in the diagnosis, treatment and prevention of conditions
CC related to diabetes, e.g. hyperglycaemia, obesity, retinopathy,
CC polynuropathy, atherosclerosis, anaemia, stroke, gangrene, impotence,
CC infection, cataract, renal disorders, or endocrine disorders. The present
CC sequence was used to illustrate the invention.
XX SQ
XX Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;

Query Match 82.9%; Score 17.4; DB 8; Length 32681;
Best Local Similarity 94.7%; Pred. No. 2.7e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTCTAG 19
Db 16526 CAGTGACAGCAGGCTCTAG 16508

RESULT 21
ADC20949/c
ID ADC20949 standard; DNA; 32681 BP.
XX AC
XX ADC20949;
XX AC
XX 18-DEC-2003 (first entry)
DT DT
XX Human secreted protein-related DNA sequence #367.
XX DE
XX Human secreted protein-related DNA sequence #367.
XX KW
XX Gene therapy; human; secreted protein; haemopoietic disorder;
KW haematological disorder; anaemia; haemophilia; inflammatory disorder;
KW inflammatory bowel disease; Crohn's disease; neoplastic disease; cancer;
KW leukaemia; wound healing; epithelial cell proliferation disorder;
KW immune disorder; autoimmune disorder; asthmatic disorder;
KW cardiovascular disorder; atherosclerosis; myocarditis;
KW infectious disease; HIV; AIDS; endocrine disorder; diabetes; ds.
KW gastrointestinal disorder; duodenal ulcer; gastroenteritis; gene; ds.
XX OS
XX Homo sapiens.
XX PN
XX WO200292787-A2.
XX PD
XX 21-NOV-2002.
XX PF
XX 26-MAR-2002; 2002WO-US009257.
XX PP

```

PR 27-MAR-2001; 2001US-0278650P.
 PR 12-SEP-2001; 2001US-00950082.
 PR 12-SEP-2001; 2001US-00950083.
 XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 PI Rosen CA, Ruben SM;
 XX
 DR WPI; 2003-129287/12.
 XX
 PT New human secreted proteins and nucleic acid molecules, useful for
 PT preparing a diagnostic or pharmaceutical composition for diagnosing,
 PT preventing or treating hematopoietic or hematologic disorders, e.g.
 PT anemia or hemophilia.
 XX
 XX Disclosure; SEQ ID NO 903; 1512pp; English.
 XX
 CC The invention comprises the amino acid and coding sequences of human
 CC secreted proteins. The DNA and protein sequences of the invention are
 CC useful for detecting, preventing, diagnosing, prognosticating, treating
 CC or ameliorating: hematopoietic or hematological disorders (e.g. anaemia
 CC and hemophilia); inflammatory disorders (e.g. inflammatory bowel disease
 CC and Crohn's disease); neoplastic disease (e.g. cancer and leukaemia);
 CC wound healing and disorders of epithelial cell proliferation; immune
 CC disorders (e.g. autoimmune disorders and asthmatic disorders);
 CC cardiovascular disorders (e.g. atherosclerosis and myocarditis);
 CC infectious disease (e.g. HIV/AIDS); endocrine disorders (e.g. diabetes);
 CC and gastrointestinal disorders (e.g. duodenal ulcers and
 CC gastroenteritis). The present DNA sequence was used in the
 CC exemplification of the invention.
 XX
 SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;
 XX
 Query Match 82.9%; Score 17.4; DB 10; Length 32681;
 Best Local Similarity 94.7%; Pred. No. 2.7e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 XX
 QY 1 CAGTGACATGCAGGCTCTAG 19
 ||||| ||||| ||||| ||||| |||||
 Db 16526 CAGTGACAGCGAGGCTCTAG 16508
 XX
 RESULT 22
 ABZ68053/C
 ID ABZ68053 standard; DNA; 32681 BP.
 XX
 AC ABZ68053;
 XX
 DT 26-MAR-2003 (first entry)
 XX
 DE Human secreted protein encoding genomic DNA SEQ ID NO 1576.
 XX
 KW Human; secreted protein; nontropic; neuroprotective; cytostatic;
 KW virucide; dermatological; immunosuppressive; antiinflammatory; anti-HIV;
 KW vulnary; antibacterial; antiparkinsonian; antiscikling; antianaemic;
 KW antiarthritic; cancer; antirheumatic; hepatotropic; cerebroprotective;
 KW antiinflammatory; anti allergic; antidiabetic; antilulcer; anticonvulsant;
 KW antifungal; antiparasitic; cardiac; immune disorder; infection; vaccine;
 KW cardiovascular disorder; neurological disease; nephrotropic;
 KW gene therapy; gene; ds.
 XX
 OS Homo sapiens.
 XX
 PN WO20027186-A2.
 XX
 PD 03-OCT-2002.
 XX
 PF 26-MAR-2002; 2002WO-US009188.
 XX
 PR 27-MAR-2001; 2001US-0278650P.
 PR 12-SEP-2001; 2001US-00950082.
 PR 12-SEP-2001; 2001US-00950083.
 XX

PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 PI Rosen CA, Ruben SM;
 XX
 DR WPI; 2003-040583/03.
 XX
 PT New human secreted proteins encoded by genes contained in cDNA clones
 PT (e.g. HGCAC19), useful for preventing, treating or diagnosing e.g. AIDS,
 PT multiple sclerosis, herpes virus, leukemia, tick-borne encephalitis or
 PT West Nile fever.
 XX
 XX Disclosure; Page 2201-2209; 2423pp; English.
 XX
 CC The invention relates to novel human genes (ABZ66891-ABZ68209) and the
 CC encoded secreted proteins (ABP99470-ABP99872) useful for preventing,
 CC treating or ameliorating medical conditions e.g. by protein or gene
 CC therapy. The genes are isolated from a range of human tissues disclosed
 CC in the specification. The nucleic acids, proteins, antibodies and
 CC (ant)agonists are useful in the diagnosis, treatment and prevention of:
 CC (a) cancer, e.g. breast and ovarian cancer and other cancers of the
 CC adrenal gland, bone, bone marrow, breast, gastrointestinal tract, liver,
 CC lung or urogenital; (b) immune disorders e.g. Addison's disease,
 CC allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis,
 CC diabetes mellitus, Crohn's disease, multiple sclerosis, rheumatoid
 CC arthritis and ulcerative colitis; (c) cardiovascular disorders such as
 CC myocardial ischaemias; (d) wound healing; (e) neurological diseases e.g.
 CC cerebral anoxia and epilepsy; and (f) infectious diseases such as viral,
 CC bacterial, fungal and parasitic infections
 XX
 SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;
 XX
 Query Match 82.9%; Score 17.4; DB 10; Length 32681;
 Best Local Similarity 94.7%; Pred. No. 2.7e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 XX
 QY 1 CAGTGACATGCAGGCTCTAG 19
 ||||| ||||| ||||| ||||| |||||
 Db 16526 CAGTGACAGCGAGGCTCTAG 16508
 XX
 RESULT 23
 ABX46275
 ID ABX46275 standard; cDNA; 320 BP.
 XX
 AC ABX46275;
 XX
 DT 21-FEB-2003 (first entry)
 XX
 DE Bovine EST associated with lactation/muscle/fat deposition #11440.
 XX
 KW Bovine; ss; EST; expressed sequence tag; lactation; LMFD;
 KW muscle deposition; fat deposition; genome mapping; gene identification;
 KW gene analysis; cattle breeding.
 XX
 OS Bos Taurus.
 XX
 PN US2002137139-A1.
 XX
 PD 26-SEP-2002.
 XX
 PF 24-SEP-2001; 2001US-00960352.
 XX
 PR 12-JAN-1999; 99US-0115707P.
 PR 11-JAN-2000; 2000US-00480902.
 XX
 PA (BYAT/) BYATT J C.
 PA (MATH/) MATHIALAGAN N.
 PA (TAON/) TAO N.
 PA (WARR/) WARREN W C.
 XX
 PI Byatt JC, Mathialagan N, Tao N, Warren WC;
 XX WPI; 2003-110599/10.
 DR

XX New nucleic acid associated with lactation, and muscle and fat
PT deposition, useful for genome mapping, gene identification and analysis,
PT cattle breeding, or for genetically improving cattle.
PS Claim 2; SEQ ID NO 11440; 245pp; English.
XX The invention relates to a purified nucleic acid molecule associated with
CC lactation or muscle and fat deposition (designated LMFD), derived from
CC cattle, and the LMFD nucleic acid can specifically hybridise to a second
CC nucleic acid molecule comprising any of 15112 nucleotide sequences,
CC appearing as ABX34836-ABX4947, or complements of them. Also included are
CC ; (1) a transformed cell having a nucleic acid comprising an LMFD nucleic
CC acid linked to a promoter and a 3' non- translated sequence that
CC functions in the cell to cause termination of transcription and addition
CC of polyadenylated ribonucleotides to a 3' end of the mRNA molecule; and
CC (2) determining a level or pattern of a molecule in a bovine cell or
CC tissue comprising: (a) incubating a marker nucleic acid (comprising any
CC of the 15112 nucleic acid sequences or its complement or fragment) with a
CC complementary nucleic acid molecule obtained from the bovine cell or
CC tissue, where hybridisation between the marker nucleic acid and the
CC complementary nucleic acid permits the detection of the molecule; and (b)
CC detecting the level or pattern of the complementary nucleic acid, where
CC the detection of the complementary nucleic acid is predictive of the
CC level or pattern of the molecule. The LMFD nucleic acid is used for
CC determining a level or pattern of a molecule in a bovine cell or tissue.
CC It is useful for genome mapping, gene identification and analysis, cattle
CC breeding, preparation of constructs for use in cattle gene expression, or
CC for genetically improving cattle. The present sequence is one of the
CC 15112 bovine LMFD EST (expressed sequence tag) nucleic acids. Note: The
CC present sequence was not shown in the specification but was obtained in
CC electronic format from the USPTO web site:
CC seqdata.uspto.gov/sequence.html?DocID=20020137139
XX
XX Sequence 320 BP; 78 A; 80 C; 97 G; 65 T; 0 U; 0 Other;
SQ
Query Match 80.0%; Score 16.8; DB 8; Length 320;
Best Local Similarity 90.0%; Pred. No. 3.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
OY 2 AGTGACATGCGAGGTCTAGCT 21
Db 144 AATGACATGCGAGGTCTACCT 163
RESULT 24
AAS00624
ID AAS00624 standard; DNA; 36221 BP.
XX AAS00624;
AC AAS00624;
XX 07-SBP-2001 (first entry)
DT Human death-associated protein 6 (DAXX) gene.
DE
XX Death-associated protein 6; DAXX; polymorphism; haplotype pair; human;
KW immune disorder; autoimmune disease; population diversity; ds;
KW paternity testing; anthropological lineage; forensic application.
XX Homo sapiens.
XX
FH Key Location/Qualifiers
FT variation replace(26869,G)
FT /*tag= a
FT variation replace(26870,T)
FT /*tag= b
FT variation replace(27145,A)
FT /*tag= c
FT variation replace(27239,G)
FT /*tag= d
FT variation replace(27620,T)
FT /*tag= e
FT variation replace(27788,G)

FT variation /*tag= f
FT replace(27806,T)
FT /*tag= g
FT variation replace(27816,T)
FT /*tag= h
FT variation replace(27869,T)
FT /*tag= i
FT variation replace(27905,A)
FT /*tag= j
FT variation replace(27916,C)
FT /*tag= k
FT variation replace(28194,T)
FT /*tag= l
FT variation replace(28339,T)
FT /*tag= m
FT variation replace(28470,C)
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FT variation replace(29010,T)
FT /*tag= o
FT variation replace(30235,T)
FT /*tag= p
FT variation replace(30665,A)
FT /*tag= q
FT variation replace(30666,T)
FT /*tag= r
FT variation replace(30752,T)
FT /*tag= s
FT variation replace(31916,T)
FT /*tag= t
XX WO200125245-A2.
XX 12-APR-2001.
XX 05-OCT-2000; 2000WO-US027487.
XX 06-OCT-1999; 99US-0157909P.
XX (GENA-) GENAISSANCE PHARM INC.
XX Chew A, Choi JY, Denton RR, Nandabalan K, Stephens JC;
XX WPI; 2001-308220/32.
XX New human death-associated protein 6 (DAXX) gene variants comprising 19
XX polymorphic sites useful in studying the effect of variation on the
XX biological activity of DAXX and in developing drugs targeting the
XX protein.
XX Claim 1; Fig 1; 97pp; English.
XX The sequence represents a DNA encoding human death-associated protein 6
XX (DAXX). This gene may comprise one or more polymorphisms at specific
XX nucleotide positions to form one of nineteen possible polymorphic
XX variants. Associations between a trait and a genotype or a haplotype of
XX the DAXX gene can be identified by comparing the frequency of the
XX genotype or haplotype in a population exhibiting the trait with that of a
XX reference population. A higher frequency in the trait population
XX indicates an association. Methods involving genotyping or haplotyping of
XX the DAXX gene of an individual can lead to prediction of haplotype pairs
XX for the DAXX gene of related individuals, and may be useful in studying
XX the expression and biological function of DAXX, as well as in developing
XX drugs targeting this protein. Polymorphic variants of DAXX are useful in
XX studying the effect of the variation on the biological activity of DAXX
XX as well as on the binding affinity of candidate drugs targeting DAXX
XX the treatment of autoimmune diseases and other immune disorders.
XX Polymorphism is also useful for studying population diversity,
XX anthropological lineage, paternity testing, forensic applications, and
XX for identifying associations between the DAXX genetic variation and a
XX trait such as level of drug response or susceptibility to disease. DAXX
XX proteins may be used to measure binding affinities of one or more
XX candidate drugs targeting the DAXX protein

SQ Sequence 36221 BP; 8897 A; 8473 C; 9437 G; 9414 T; 0 U; 0 Other;
 Query Match 80.0%; Score 16.8; DB 4; Length 36221;
 Best Local Similarity 90.0%; Pred. No. 5.2e+02;
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 AGTGACATGCAGGTCTAGCT 21
 ||||| ||||| ||||| |||||
 Db 21390 AGTGACAGGCGAGTCTAGCT 21409

RESULT 25
 ACN45158
 ID ACN45158 standard; DNA; 72705 BP.
 XX
 AC ACN45158;
 XX
 DT 18-NOV-2004 (first entry)
 XX
 DE Human genomic sequence hCG25130.
 XX
 KW Cytostatic; carcinoma; lymphoma; cancer; human; gene; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO2003073826-A2.
 XX
 PD 12-SEP-2003.
 XX
 PF 28-FEB-2003; 2003WO-US006235.
 XX
 PR 01-MAR-2002; 2002US-00087192.
 XX
 PA (SAGR-) SAGRES DISCOVERY.
 XX
 PI Morris DW;
 XX
 DR WPI; 2003-328604/31.
 XX
 PT Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
 PT comprises a nucleotide sequence.
 XX
 PS Claim 1; SEQ ID NO 1966; Opp; English.
 XX
 CC The present invention relates to novel DNA and protein sequences which
 CC are associated with carcinomas. The sequences are useful for: (i) for
 CC screening drug candidates; (ii) for screening of bioactive agent capable
 CC of binding to Carcinoma Associated Protein (CAP); (iii) for screening of
 CC a bioactive agent capable of modulating the activity of CAP; (iv) for
 CC evaluating the effect of a candidate carcinoma drug; (v) for diagnosing
 CC carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating
 CC carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip;
 CC (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
 CC determining Carcinoma Associated (CA) gene copy number. In addition, the
 CC CA genes are useful as DNA vaccines and the CAP are useful as markers of
 CC carcinoma including lymphoma. The present sequence is one such CA coding
 CC sequence. Note: This patent is an equivalent to basic patent
 CC US2002182586A1, for which no sequence data was published
 XX

SQ Sequence 72705 BP; 18277 A; 18952 C; 18052 G; 17424 T; 0 U; 0 Other;
 Query Match 78.1%; Score 16.4; DB 11; Length 72705;
 Best Local Similarity 94.4%; Pred. No. 8.7e+02;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 AGTGACATGCAGGTCTAG 19
 ||||| ||||| ||||| |||||
 Db 32065 AGTGACATGCAGGTCTAG 32082

Search completed: September 6, 2005, 20:39:33
 Job time : 194.656 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 17:45:55 ; Search time 1572.31 Seconds
(without alignments)
532.600 Million cell updates/sec

Title: US-10-729-421-35

Perfect score: 22

Sequence: 1 agccctcttcagtcacatcaag 22

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 100 summaries

Database :

EST:**

1: gb_est1:**

2: gb_est2:**

3: gb_hic:**

4: gb_est3:**

5: gb_est4:**

6: gb_est5:**

7: gb_est6:**

8: gb_gsal:**

9: gb_gse2:**

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	19	86.4	1174	8	CC229626 CH261-46H
2	18.8	85.5	767	2	BE541043 601064391
3	18.8	85.5	872	2	BF687801 602066853
C 4	18.4	83.6	935	9	CL901625 CSHC1000
C 5	17.8	80.9	169	1	AA693379 ah21906.s
C 6	17.8	80.9	185	2	BF833890 RC1-HT088
7	17.8	80.9	193	2	AW800621 NR1-UM006
8	17.8	80.9	205	2	BF111499 7128f04.x
C 9	17.8	80.9	270	1	AA884812 am28805.s
C 10	17.8	80.9	270	2	AW819544 RCS-ST029
C 11	17.8	80.9	271	1	AA437133 zv53e10.s
C 12	17.8	80.9	271	2	AA467041 ha08c03.x
13	17.8	80.9	289	2	BB720494 BB720494
14	17.8	80.9	285	4	BM151302 TCBAF1D61
C 15	17.8	80.9	309	1	AI203209 gr23h08.x
16	17.8	80.9	310	7	T46996 yb12b06.s1
17	17.8	80.9	328	1	AA969466 co81d01.s
C 18	17.8	80.9	328	2	AW615159 hg73h02.x
C 19	17.8	80.9	339	1	AI307235 tb18c07.x
C 20	17.8	80.9	339	1	AI631503 wa89909.x
C 21	17.8	80.9	339	1	AI633750 tt28b04.x
C 22	17.8	80.9	339	1	AI954798 wq33d07.x
C 23	17.8	80.9	339	1	AI954834 wq33h04.x
C 24	17.8	80.9	339	1	AI962281 wq46e04.x

T46995 yb12b06.r1	339	7	T46995	80.9	17.8	C 25
AW291932 UI-H-BI2-	340	2	AW291932	80.9	17.8	C 26
AW237469 xm72c11.x	345	2	AW237469	80.9	17.8	C 27
AZ693999 AST-1HDB2	350	8	AZ693999	80.9	17.8	C 28
AT749323 nyl2c01.s	356	1	AT749323	80.9	17.8	C 29
AI150471 qf14d04.x	376	1	AI150471	80.9	17.8	C 30
AA535707 nf88d04.s	389	1	AA535707	80.9	17.8	C 31
AA535707 nf88d04.s	389	1	AA535707	80.9	17.8	C 32
AA291203 UI-H-BI2-	398	1	AA291203	80.9	17.8	C 33
AI242291 q116g02.x	404	1	AI242291	80.9	17.8	C 34
AI074659 ox82g07.s	416	1	AI074659	80.9	17.8	C 35
CN545186 EST 17130	423	7	CN545186	80.9	17.8	C 36
AI041258 ov66a02.x	428	1	AI041258	80.9	17.8	C 37
BM998692 UI-H-DT1-	435	5	BM998692	80.9	17.8	C 38
AA417921 zv94b03.s	444	1	AA417921	80.9	17.8	C 39
AA834127 of26g07.s	447	1	AA834127	80.9	17.8	C 40
AW073711 x501h10.x	447	2	AW073711	80.9	17.8	C 41
AI148470 t94he05.x	448	1	AI148470	80.9	17.8	C 42
AA768438 ob22f08.s	450	1	AA768438	80.9	17.8	C 43
AA418172 zv94f03.f	450	1	AA418172	80.9	17.8	C 44
BQ942517 AGENCOURT	452	5	BQ942517	80.9	17.8	C 45
BP305540 BP305540	455	5	BP305540	80.9	17.8	C 46
BY560157 BY560157	457	6	BY560157	80.9	17.8	C 47
BE379976 601159478	460	2	BE379976	80.9	17.8	C 48
AA159601 z080a04.s	463	1	AA159601	80.9	17.8	C 49
AW275079 xm82c01.x	463	2	AW275079	80.9	17.8	C 50
BQ012758 UI-1-BC1P	470	5	BQ012758	80.9	17.8	C 51
AI088652 qb14a07.x	484	1	AI088652	80.9	17.8	C 52
AA936255 on75b04.s	490	1	AA936255	80.9	17.8	C 53
AW471206 xv13e09.x	494	2	AW471206	80.9	17.8	C 54
BX279662 BX279662	506	5	BX279662	80.9	17.8	C 55
BQ774467 UI-H-EZ1-	516	5	BQ774467	80.9	17.8	C 56
BE440143 HTM1-954R	526	2	BE440143	80.9	17.8	C 57
AI400162 t967g09.x	530	1	AI400162	80.9	17.8	C 58
AW128873 x889c02.x	534	2	AW128873	80.9	17.8	C 59
AA864874 oh03d08.s	541	1	AA864874	80.9	17.8	C 60
BF890876 PM2-MT010	546	2	BF890876	80.9	17.8	C 61
BI962926 i660d12.y	547	4	BI962926	80.9	17.8	C 62
BI966885 i662f08.x	549	4	BI966885	80.9	17.8	C 63
BI963087 i662f08.y	550	4	BI963087	80.9	17.8	C 64
AI088606 qb14e01.x	551	1	AI088606	80.9	17.8	C 65
AV735110 AV735110	555	1	AV735110	80.9	17.8	C 66
AW970416 EST382497	558	2	AW970416	80.9	17.8	C 67
CB267636 1006542.H	563	6	CB267636	80.9	17.8	C 68
AV715652 AV715652	568	1	AV715652	80.9	17.8	C 69
BF970233 602273470	568	4	BF970233	80.9	17.8	C 70
BG723880 602697376	568	4	BG723880	80.9	17.8	C 71
AL701719 DKF2P686H	569	1	AL701719	80.9	17.8	C 72
BX094624 BX094624	569	5	BX094624	80.9	17.8	C 73
BM670715 UI-E-DX1-	576	4	BM670715	80.9	17.8	C 74
AI309768 q075b03.x	577	1	AI309768	80.9	17.8	C 75
BF379530 BF379530	582	5	BF379530	80.9	17.8	C 76
BI966756 i660d12.x	584	4	BI966756	80.9	17.8	C 77
CB160326 K-EST0220	585	6	CB160326	80.9	17.8	C 78
BE739363 601556586	593	2	BE739363	80.9	17.8	C 79
AV716017 AV716017	613	1	AV716017	80.9	17.8	C 80
CV027350 5580.FULL1	618	7	CV027350	80.9	17.8	C 81
BG500991 602546583	620	4	BG500991	80.9	17.8	C 82
BE739605 601556586	623	2	BE739605	80.9	17.8	C 83
BI553318 603193449	643	4	BI553318	80.9	17.8	C 84
BG535175 602562794	647	4	BG535175	80.9	17.8	C 85
BG505886 601859895	654	4	BG505886	80.9	17.8	C 86
BM713190 UI-E-EJ0-	657	4	BM713190	80.9	17.8	C 87
B63640 RPC111-2J13	657	8	B63640	80.9	17.8	C 88
BF213738 601847628	663	2	BF213738	80.9	17.8	C 89
CD639326 AGENCOURT	663	6	CD639326	80.9	17.8	C 90
CE750532 t1gr-g88-	673	9	CE750532	80.9	17.8	C 91
CD641555 AGENCOURT	682	6	CD641555	80.9	17.8	C 92
AV762336 AV762336	721	1	AV762336	80.9	17.8	C 93
BE603221 HVSMH010	732	1	BE603221	80.9	17.8	C 94
AV758775 AV758775	742	1	AV758775	80.9	17.8	C 95
BI562028 603255083	765	4	BI562028	80.9	17.8	C 96
BE566608 601340137	766	2	BE566608	80.9	17.8	C 97

98 17.8 80.9 766 9 BX204503 BX204503 Danio rer
 99 17.8 80.9 775 2 BF529524 602043291
 c 100 17.8 80.9 785 1 AV757859 AV757859 AV757859

ALIGNMENTS

CC229626 1174 bp DNA linear GSS 12-MAY-2003
 CH261-46H2_Sp6.1 CH261 Gallus gallus genomic clone CH261-46H2,
 genomic survey sequence.

ACCESSION CC229626
 VERSION CC229626
 SOURCE GSS.

ORGANISM Gallus gallus (chicken)
 Gallus gallus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
 Phasianinae; Gallus.

REFERENCE 1 (bases 1 to 1174)
 Kremitzki, C., Higginbotham, J., Wylie, K., Carter, J., McPherson, J.,
 Warren, W., Graves, T., Mardis, E. and Wilson, R.
 Gallus gallus BAC End Reads

TITLE Unpublished (2003)
 JOURNAL Contact: Richard K. Wilson
 COMMENT Genome Sequencing Center
 Washington University School of Medicine
 Email: submissions@watson.wustl.edu

Insert Length: 182000 Std Error: 0.00
 Seq primer: Sp6 ATTATAGTGACACTATAG
 Class: BAC ends

High quality sequence start: 329
 High quality sequence stop: 482.

FEATURES Location/Qualifiers

source 1..1174

/organism="Gallus gallus"

/mol_type="genomic DNA"

/strain="Red Jungle Fowl"

/db_xref="taxon:9031"

/clone="CH261-46H2"

/sex="female"

/cell_line="UCD001, inbred 256"

/clone_lib="CH261"

/notes="Vector: pTARBAC2.1; Site 1: EcoRI; Site 2: EcoRI;
 CH261 Female Chicken library - for library and clone
 ordering information: http://www.chori.org/bacpac"

ORIGIN

Query Match 86.4%; Score 19; DB 8; Length 1174;
 Best Local Similarity 100.0%; Pred. No. 3.8e+02;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 CCTCTTCAGTCCCAATCAAG 22

|||||

Db 399 CCTCTTCAGTCCCAATCAAG 417

|||||

RESULT 2

BE541043

LOCUS 601064391F1 NIH_MGC_10 Homo sapiens cDNA clone IMAGE:3450647 5',
 mRNA sequence.

DEFINITION BE541043

ACCESSION BE541043

VERSION BE541043

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 767)

AUTHORS NIH-MGC http://mgs.nci.nih.gov/.

TITLE JOURNAL
 COMMENT

National Institutes of Health, Mammalian Gene Collection (MGC)
 Unpublished (1999)
 Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: ATCC

cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: Incyte Genomics, Inc.
 DNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov

Plate: LLAM8429 row: f column: 24

High quality sequence stop: 523.

FEATURES Location/Qualifiers

source 1..767

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:3450647"

/cell_line="MGC36"

/lab_host="DH10B"

/clone_lib="NIH_MGC_10"

/note="Organ: cervix; Vector: pCMV-SPORT6; Site 1: NotI;
 Site 2: SalI; Cloned unidirectionally. Primer: Oligo dt.
 Average insert size 1.5 kb. Library prepared by Life
 Technologies."

ORIGIN

Query Match 85.5%; Score 18.8; DB 2; Length 767;
 Best Local Similarity 90.9%; Pred. No. 4.5e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCCAATCAAG 22

|||||

Db 107 AGTCCTCTTCAGTCCCAATCAAG 128

|||||

RESULT 3

BF687801

LOCUS 602066853F1 NIH_MGC_57 Homo sapiens cDNA clone IMAGE:4065901 5',
 mRNA sequence.

DEFINITION BF687801

ACCESSION BF687801

VERSION BF687801

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 872)

AUTHORS NIH-MGC http://mgs.nci.nih.gov/.

TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

JOURNAL Unpublished (1999)

COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: ATCC

cDNA Library Preparation: CLONETECH Laboratories, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov

Plate: LLCM902 row: j column: 14

High quality sequence stop: 623.

FEATURES Location/Qualifiers

source 1..872

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:4065901"

/tissue_type="glioblastoma"

/lab_host="DH10B (T1 phage-resistant)"

/clone_lib="NIH_MGC_57"

ORIGIN

Query Match 86.4%; Score 19; DB 8; Length 1174;
 Best Local Similarity 100.0%; Pred. No. 3.8e+02;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 CCTCTTCAGTCCCAATCAAG 22

|||||

Db 399 CCTCTTCAGTCCCAATCAAG 417

|||||

RESULT 2

BE541043

LOCUS 601064391F1 NIH_MGC_10 Homo sapiens cDNA clone IMAGE:3450647 5',
 mRNA sequence.

DEFINITION BE541043

ACCESSION BE541043

VERSION BE541043

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 767)

AUTHORS NIH-MGC http://mgs.nci.nih.gov/.

/note="Organ: brain; Vector: pDNR-LIB (Clontech); Site:1: SfiI (ggcgctcgcc); Site 2: SfiI (ggcattatggcc); Double-stranded cDNA was prepared from cell line RNA. 5' and 3' adaptors were used in cloning as follows: 5' adaptor sequence: 5'-CACGCCATTATGCG-3' and 3' adaptor sequence: 5'-ATTCTAGAGCGCGCGCCGACATG-dT(30)BN-3' (where B = A, C, or G and N = A, C, G, or T). Average insert size 1.55 kb (range 0.9-4.0 kb). 12/15 colonies contained inserts by PCR. This library was enriched for full-length clones and was constructed by Clontech Laboratories (Palo Alto, CA)."

ORIGIN

Query Match 85.5%; Score 18.8; DB 2; Length 872;
Best Local Similarity 90.9%; Pred. No. 4.5e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCAAG 22
||| ||||| ||||| ||||| |||||
Db 319 AGTCTCTTCAGTCCAATCAAG 340

RESULT 4
CL901625/c

LOCUS CSHCI000 1639HC library Triticum aestivum genomic clone
DEFINITION 1639HC06E05, genomic survey sequence.

ACCESSION CL901625
VERSION CL901625.1 GI:51663670
KEYWORDS GSS.

SOURCE Triticum aestivum (bread wheat)
ORGANISM Triticum aestivum

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Poidea; Triticeae; Triticum.

REFERENCE 1 (bases 1 to 935)

AUTHORS Lamoureux, D., Peterson, D.G., Li, W., Fellers, J.P. and Gill, B.S.
TITLE Cot-based cloning and sequencing (CBCS) efficiently removes sequence repeats and increases gene ratio in bread wheat
JOURNAL Unpublished (2004)

COMMENT Contact: Gill BS

Department of Plant Pathology
Kansas State University
4024 Throckmorton, Manhattan, KS 66506-5502, USA
Tel: 785 532 1391
Fax: 785 532 5692
Email: begill@ksu.edu
Seq primer: T7
Class: sheared ends.

FEATURES

Location/Qualifiers
1..935
/organism="Triticum aestivum"
/mol_type="genomic DNA"
/cultivar="Chinese Spring"
/db_xref="taxon:4565"
/clone="1639HC06E05"
/tissue_type="whole plant"
/dev_stage="young shoot"
/clone_lib="1639HC library"

ORIGIN

Query Match 83.6%; Score 18.4; DB 9; Length 935;
Best Local Similarity 95.0%; Pred. No. 7.1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 CCCTCTTCAGTCCAATCAAG 22
||| ||||| ||||| ||||| |||||
Db 216 CCCTCTTCAGTCCAATCAAG 197

RESULT 5

AA693379/c
LOCUS

DEFINITION

ah21g06.s1 Soares parathyroid_tumor NHPA Homo sapiens cDNA clone

1239514 3' similar to TR:Q13227 Q13227 GPS2. ;, mRNA sequence.

ACCESSION AA693379

VERSION AA693379.1 GI:2694317

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 169)

AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

COMMENT Contact: Robert Strausberg, Ph.D.

Email: cgapbs@mail.nih.gov

CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima

Bonaldo, Ph.D.

CDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LINL at:

www.bio.lnl.gov/bbrp/image/image.html

Trace considered overall poor quality

Possible reversed clone; similarity on wrong strand

Insert Length: 912 Std Error: 0.00

Seq primer: -40ml3 fwd. ET from Amersham

High quality sequence stop: 1.

Location/Qualifiers

1..169

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="1239514"

/tissue_type="parathyroid tumor"

/dev_stage="adult"

/lab_host="DH10B (ampicillin resistant)"

/clone_lib="Soares parathyroid tumor NHPA"

/note="Organ: parathyroid gland; Vector: pT7T3D

(Pharmacia) with a modified polylinker; Site 1: Not I;

Site 2: Eco RI; 1st strand cDNA was primed with a Not I -

oligo(dT) primer

[5'-TGTTACCAATCTGAAGTGGAGCGCGCCACCAATTTTTTTTTTTTTTTT

TTTTT-3'], double-stranded cDNA was size selected, ligated

to Eco RI adapters (Pharmacia), digested with Not I and

cloned into the Not I and Eco RI sites of a modified pT7T3

vector (Pharmacia). Library went through one round of

normalization to a Cot = 5. Library constructed by Bento

Soares and M.Fatima Bonaldo. RNA from sporadic parathyroid

adenomas was kindly provided by Dr. Stephen Marx, National

Institute of Diabetes and Digestive and Kidney Diseases,

NIH."

ORIGIN

Query Match 80.9%; Score 17.8; DB 1; Length 169;
Best Local Similarity 90.5%; Pred. No. 1.1e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GCCCTCTTCAGTCCAATCAAG 22

||| ||||| ||||| ||||| |||||

Db 130 GCCATCTTCAGCCCAATCAAG 110

RESULT 6

BF833890/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

BF833890 185 bp mRNA linear EST 13-JAN-2001
RC1-HT0881-041100-019-b04 HT0881 Homo sapiens cDNA, mRNA sequence.

BF833890

BF833890.1 GI:12183723

EST.

Homo sapiens (human)

Homo sapiens

Homo sapiens

TITLE RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura,T., et al. 2001)

JOURNAL Unpublished (2001)

COMMENT Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-resgsc.riken.jp, URL: http://genome.gsc.riken.jp/
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)
wagi, K., Fujiwaki, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)
Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara, Y. and Hayashizaki, Y.
Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
Please visit our web site (http://genome.gsc.riken.go.jp) for further details.

FEATURES

source

1. .289
Location/Qualifiers
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="C730036G02"
/sex="male"
/tissue_type="liver tumor"
/dev_stage="adult"
/lab_host="DH108"
/clone_lib="RIKEN full-length enriched, adult male liver tumor"

/notes="Site 1: SalI; Site 2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5'
GAGAGAGAGCGCGCACTCGAGTTTTTTTTTTT 3']. cDNA was prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. Second strand cDNA was prepared with the primer adapter of sequence [5'
GAGAGAGATTTCGAGTTAATTAATTAATCCCCCCCCC 3']. cDNA was cleaved with BamHI and XhoI. Vector: a modified pluescript KS(+) after bulk excision from Lambda FIC I. Tissue was provided by William A. Held, Roswell Park Cancer Institute, Department of Molecular and Cellular Biology, Elm and Carlton Streets, Buffalo, NY 14263, whose assistance we gratefully acknowledge."

ORIGIN

Query Match 80.9%; Score 17.8; DB 2; Length 289;
Best Local Similarity 90.5%; Pred. No. 1.2e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAA 21
|||||||
DB 10 AGCCCTCTTCATTCCTCAAC 30
|||||||

RESULT 14

BM151302
LOCUS
DEFINITION

ACCESSION
VERSION
KEYWORDS
SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

source

1. .295
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="TCBAP6134"
/sex="male"
/tissue_type="leukopheresis"
/cell_type="pre-B cell"
/dev_stage="pediatric 2 years"
/lab_host="DH108"
/clone_lib="pediatric pre-B cell acute lymphoblastic leukemia Baylor-HGSC project=TCBA"
/note="Vector: lambda pSB, Site 1: BamHI; Site 2: EcoRI; First strand cDNA was primed with an anchored XhoI-oligo(dT) primer [5'GGAGACTCGAGCGCGCAGGAG(T)VN 3'; V=A,C,G; N=A,C,G,T] and then dg tailed. Second strand was primed with a BamHI-dC primer [5'AGAGACTCGGATCCGCGCGCAATTAATAAT(C) 3']. Double-stranded cDNA was then digested with BamHI and XhoI and directionally cloned into the BamHI and SalI sites of lambda pSB vector. Library was constructed by Wei Yu at RIKEN normalization. Library was constructed by Wei Yu at RIKEN of Japan (Carninci P, Westover A, Nishiyama Y, Ohsumi T, Itoh M, Nagaoka S, Sasaki, Okazaki Y, Muramatsu M, Schneider C, Hayashizaki Y, High efficiency selection of full-length cDNA by improved biotinylated cap trapper., DNA Res 4: 1, 61-6, Feb 28, 1997)"

ORIGIN

Query Match 80.9%; Score 17.8; DB 4; Length 295;
Best Local Similarity 90.5%; Pred. No. 1.2e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GCCCTCTTCAGTCCCAATCAAG 22
|||||||
DB 269 GCCATCTTCAGTCCCAATCCAG 289
|||||||

RESULT 15

AI203209/c
LOCUS
DEFINITION

ACCESSION
VERSION
KEYWORDS

AI203209
qr23h08.x1 NCI_CGAP_GC6 Homo sapiens cDNA clone IMAGE:1941759 3', mRNA sequence.

AI203209

AI203209

AI203209.1 'GI:3755815

EST.

BM151302 295 bp mRNA linear EST 30-NOV-2001
TCBAP106134 Pediatric pre-B cell acute lymphoblastic leukemia
Baylor-HGSC project=TCBA Homo sapiens cDNA clone TCBAP6134, mRNA sequence.

BM151302
EST.

BM151302.1 GI:17174603

EST.

Homo sapiens (human)

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 295)

Wei, Y., Tsang, Y. T. M., Mei, G., Ku, J. M., Ali-Osman, F. R. Jr., Gunaratne, P. H., Muzny, D., Bouck, J., Gibbs, R. A. and Margolin, J. F.

Pediatric Leukemia cDNA Sequencing Project (2001)

Unpublished (2001)

Contact: Dr. Judith F. Margolin

Texas Children's Cancer Center and Human Genome Sequencing Center

at Baylor College of Medicine

1102 Bates MC3-3320 Houston, TX 77030, USA

Tel: 832-824-4536

Fax: 832-825-4038

Email: clones@tccc.org

Seq primer: M13 primer.

Trace considered overall poor quality
 Insert Length: 1094 Std Error: 0.00
 Seq primer: -40m13 fwd. ET from Amersham
 High quality sequence stop: 1.

FEATURES

source
 1. .328
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:1572577"
 /tissue_type="2 pooled tumors (clear cell type)"
 /lab_host="DH10B"
 /clone_lib="NCI CGAP Kids"
 /notes="Organ: kidney; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' AACTGGAGAAATTCGCGCGCAATATTTTTTTTTTTT 3'], double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo. "

ORIGIN

Query Match 80.9%; Score 17.8; DB 1; Length 328;
 Best Local Similarity 90.5%; Pred. No. 1.2e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GCCCTCTTCAGTCCAATCAAG 22

Db 153 GCCATCTTCAGTCCAATCCAG 173

RESULT 18

AW615159/c
 LOCUS hg73h02.x1 NCI_CGAP_GC6 Homo sapiens cDNA linear EST 23-MAR-2000
 DEFINITION mRNA sequence.
 ACCESSION AW615159 328 bp mRNA
 VERSION AW615159.1 GI:7320345
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 328)
 AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index

JOURNAL

COMMENT
 Unpublished (1997)
 Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo, Ph.D.

cDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov/image/html/iresources.shtml
 Seq primer: -40UP from Gibco.

FEATURES

source
 1. .328
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:2951283"
 /tissue_type="pooled germ cell tumors"
 /lab_host="DH10B"
 /clone_lib="NCI_CGAP_GC6"
 /notes="Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; Plasmid DNA

ORIGIN

Query Match 80.9%; Score 17.8; DB 2; Length 328;
 Best Local Similarity 90.5%; Pred. No. 1.2e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GCCCTCTTCAGTCCAATCAAG 22

Db 230 GCTGTTCTTCAGTCCAATCAAG 210

RESULT 19

AI307235
 LOCUS tb18c07.x1 NCI_CGAP_Kid12 Homo sapiens cDNA linear EST 08-APR-1999
 DEFINITION similar to TR:Q92478 Q92478 C-TYPE LECTIN.; mRNA sequence.

ACCESSION AI307235.1 GI:4001991

VERSION AI307235

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 339)

AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index

UNPUBLISHED (1997)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

DNA Sequencing by: Greg Lennon, Ph.D.

Clone distribution: Washington University Genome Sequencing Center

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 603 Std Error: 0.00

Seq primer: -40UP from Gibco.

Location/Qualifiers

1. .339

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:2054700"

/tissue_type="2 pooled tumors (clear cell type)"

/lab_host="DH10B"

/clone_lib="NCI_CGAP_Kid12"

/notes="Organ: Kidney; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; Plasmid DNA from the normalized library NCI_CGAP Kids was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (cloneIDs 1323912-1325831, 1471368-1472903 and 1492104-1493255). Subtraction by Bento Soares and M. Fatima Bonaldo. "

ORIGIN

Query Match 80.9%; Score 17.8; DB 1; Length 339;
 Best Local Similarity 90.5%; Pred. No. 1.2e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GCCCTCTTCAGTCCAATCAAG 22

from the normalized library NCI_CGAP GC4 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (cloneIDs 1257096-1258631, 1469064-1470983, and 1475592-1476743). Subtraction by Bento Soares and M. Fatima Bonaldo. "

```

|||||
254 GCCATCTTCAGTCCAATCAAG 274

RESULT 20
AI631503/c
LOCUS
DEFINITION
wa89g09.x1 NCI_CGAP_GC6 Homo sapiens cDNA clone IMAGE:2303392 3',
mRNA sequence.
ACCESSION
AI631503
VERSION
AI631503.1 GI:4682833
KEYWORDS
EST.
SOURCE
Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 339)
AUTHORS
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TUMOR
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL
Unpublished (1997)
COMMENT
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
R. Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Insert Length: 424 Std Error: 0.00
Seq primer: -40UP from Gibco.
Location/Qualifiers
1..339
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2303392"
/tissue_type="pooled germ cell tumors"
/lab_host="DH10B"
/clone_lib="NCI_CGAP GC6"
/notes="Vector: pT7T3D-Pac (Pharmacia) with a modified
polylinker; Site 1: Not I; Site 2: Eco RI; Plasmid DNA
from the normalized library NCI_CGAP_GC4 was prepared, and
ss circles were made in vitro. Following HAP purification,
this DNA was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from a pool
of 5,000 clones made from the same library (clonesIDs
1257096-1258631, 1469064-1470983, and 1475592-1476743).
Subtraction by Bento Soares and M. Fatima Bonaldo. "

FEATURES
source
1..339
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2303392"
/tissue_type="pooled germ cell tumors"
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ss circles were made in vitro. Following HAP purification,
this DNA was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from a pool
of 5,000 clones made from the same library (clonesIDs
1257096-1258631, 1469064-1470983, and 1475592-1476743).
Subtraction by Bento Soares and M. Fatima Bonaldo. "

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Best Local Similarity 90.5%; Pred. No. 1.2e+03;
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ACCESSION
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VERSION
AI633750.1 GI:4685080
KEYWORDS
EST.
SOURCE
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 339)
AUTHORS
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TUMOR
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL
Unpublished (1997)
COMMENT
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
R. Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
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from the normalized library NCI_CGAP_GC4 was prepared, and
ss circles were made in vitro. Following HAP purification,
this DNA was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from a pool
of 5,000 clones made from the same library (clonesIDs
1257096-1258631, 1469064-1470983, and 1475592-1476743).
Subtraction by Bento Soares and M. Fatima Bonaldo. "

ORIGIN
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ACCESSION
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VERSION
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EST.
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ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 339)
AUTHORS
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TUMOR
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL
Unpublished (1997)
COMMENT
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
R. Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.

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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
R. Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
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ss circles were made in vitro. Following HAP purification,
this DNA was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from a pool
of 5,000 clones made from the same library (clonesIDs
1257096-1258631, 1469064-1470983, and 1475592-1476743).
Subtraction by Bento Soares and M. Fatima Bonaldo. "

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Query Match 80.9%; Score 17.8; DB 1; Length 339;
Best Local Similarity 90.5%; Pred. No. 1.2e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GCCCTCTTCAGTCCAATCAAG 22
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Db 244 GCTGCTTCAGTCCAATCAAG 224

RESULT 22
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VERSION
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KEYWORDS
EST.
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ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
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AUTHORS
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TUMOR
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL
Unpublished (1997)
COMMENT
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
R. Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.

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Job time : 1578.31 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 19:12:21 ; Search time 62.9062 Seconds
(without alignments)
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Title: US-10-729-421-35

Perfect score: 22

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Listing first 100 summaries

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Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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ALIGNMENTS

RESULT 1
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; Sequence 13311, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961

; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 13311
; LENGTH: 247
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-513-999C-13311

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; Sequence 15744, Application US/09621976
; Patent No. 6639063
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Jobert, S. J.Y.

; TITLE OF INVENTION: ESTs and Encoded Human Proteins.

; FILE REFERENCE: GENSET.054PR2
; CURRENT APPLICATION NUMBER: US/09/621,976
; CURRENT FILING DATE: 2000-07-21
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; Sequence 5, Application US/09944807
; Patent No. 6773895
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; APPLICANT: Boehringer Ingelheim Pharma KG
; TITLE OF INVENTION: Method for identifying substances which positively

; TITLE OF INVENTION: influence inflammatory conditions of chronic
; FILE REFERENCE: 082_00n
; CURRENT APPLICATION NUMBER: US/09/944,807
; CURRENT FILING DATE: 2001-08-31
; PRIOR APPLICATION NUMBER: UK 0021484.1
; PRIOR FILING DATE: 2000-09-01
; NUMBER OF SEQ ID NOS: 24
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US-09-944-807-5

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RESULT 4

US-08-916-421B-1
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; Patent No. 6503729
; GENERAL INFORMATION:
; APPLICANT: Bult et al.

; TITLE OF INVENTION: Complete Genome Sequence of the Methanogenic Archaeon, Methanococcus

; Patent No. 6503729

; TITLE OF INVENTION: jannaschii

; FILE REFERENCE: PB275

; CURRENT APPLICATION NUMBER: US/08/916,421B

; CURRENT FILING DATE: 1997-08-22

; PRIOR APPLICATION NUMBER: US 60/024,428

; PRIOR FILING DATE: 1996-08-22

; NUMBER OF SEQ ID NOS: 3

; SOFTWARE: PatentIn version 3.1

; SEQ ID NO 1

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; TYPE: DNA

; ORGANISM: Methanococcus jannaschii

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; NAME/KEY: misc feature
; LOCATION: (657081)..(657081)
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; NAME/KEY: misc feature
; LOCATION: (657203)..(657203)
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; NAME/KEY: misc feature
; LOCATION: (674435)..(674435)
; OTHER INFORMATION: n equals a, t, c, or g
; NAME/KEY: misc feature
; LOCATION: (682442)..(682442)
; OTHER INFORMATION: n equals a, t, c, or g
; NAME/KEY: misc feature
; LOCATION: (713652)..(713652)
; OTHER INFORMATION: n equals a, t, c, or g
; NAME/KEY: misc feature
; LOCATION: (741684)..(741684)

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; OTHER INFORMATION: n equals a, t, c, or g
; NAME/KEY: misc feature
; LOCATION: (779455)..(779455)
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; NAME/KEY: misc feature
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; OTHER INFORMATION: n equals a, t, c, or g
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; LOCATION: (855539)..(855539)
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; LOCATION: (871619)..(871619)
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; LOCATION: (1084830)..(1084830)
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; OTHER INFORMATION: n equals a, t, c, or g
; NAME/KEY: misc feature
; LOCATION: (1119881)..(1119881)
; OTHER INFORMATION: n equals a, t, c, or g
; NAME/KEY: misc feature
; LOCATION: (1130881)..(1130881)
; OTHER INFORMATION: n equals a, t, c, or g
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; LOCATION: (1310988)..(1310988)
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; NAME/KEY: misc feature
; LOCATION: (1313224)..(1313224)
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; NAME/KEY: misc feature
; LOCATION: (1349473)..(1349473)
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; NAME/KEY: misc feature
; LOCATION: (1349491)..(1349491)
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; LOCATION: (1603734)..(1603734)
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; OTHER INFORMATION: n equals a, t, c, or g
; US-08-916-421B-1

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Query Match 80.9%; Score 17.8; DB 4; Length 1664976;
 Best Local Similarity 90.5%; Pred. No. 1.4e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAA 21
 |||||
 Db 1438420 AGCCATCTTCAGTCCCTATCAA 1438440

RESULT 5
 US-09-692-570-1
 ; Sequence 1, Application US/09692570
 ; Patent No. 6797466
 ; GENERAL INFORMATION:
 ; APPLICANT: Bult et al.
 ; TITLE OF INVENTION: Complete Genome Sequence of the Methanogenic Archaeon, Methanococcus

```
; Patent No. 6797466
; TITLE OF INVENTION: jannaschii
; FILE REFERENCE: PB275C1
; CURRENT APPLICATION NUMBER: US/09/692,570
; CURRENT FILING DATE: 2003-01-14
; PRIOR APPLICATION NUMBER: US 60/024,428
; PRIOR FILING DATE: 1996-08-22
; PRIOR APPLICATION NUMBER: US 08/916,421
; PRIOR FILING DATE: 1997-08-22
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
; LENGTH: 1664976
; TYPE: DNA
; ORGANISM: Methanococcus jannaschii
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; NAME/KEY: misc_feature
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; NAME/KEY: misc_feature
; LOCATION: (234814)..(234814)
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; OTHER INFORMATION: n equals a, t, c, or g
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; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (622708)..(622708)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (657081)..(657081)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (657203)..(657203)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (674435)..(674435)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc_feature
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; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (741684)..(741684)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
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; OTHER INFORMATION: n equals a, t, c, or g
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; NAME/KEY: misc feature
; LOCATION: (779676)..(779676)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (855539)..(855539)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (871619)..(871619)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1084830)..(1084830)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1096846)..(1096846)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1119881)..(1119881)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1130881)..(1130881)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1310988)..(1310988)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1313224)..(1313224)
; OTHER INFORMATION: n equals a, t, c, or g
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1349473)..(1349473)
; OTHER INFORMATION: n equals a, t, c, or g

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Query Match 80.9%; Score 17.8; DB 4; Length 1664976;
Best Local Similarity 90.5%; Pred. No. 1.4e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Qy 1 AGCCCTCTTCAGTCCCAATCAA 21
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Db 1438420 AGCCATCTTCAGTCCATCAA 1438440

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RESULT 6
US-08-961-083-71/c
; Sequence 71, Application US/08961083
; Patent No. 6159469
; GENERAL INFORMATION:
; APPLICANT: Choi et. al.
; TITLE OF INVENTION: Streptococcus pneumoniae Antigens and Vaccines
; NUMBER OF SEQUENCES: 452
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: Maryland
; COUNTRY: USA

```

```

; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage
; OPERATING SYSTEM: HP Vectra 486/33
; SOFTWARE: MSDOS version 6.2
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/961,083
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Brookes, A. Anders
; REGISTRATION NUMBER: 36,373
; REFERENCE/DOCKET NUMBER: PB340P2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (301) 309-8504
; TELEFAX: (301) 309-8512
; INFORMATION FOR SEQ ID NO: 71:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1855 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; US-08-961-083-71

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Query Match 78.2%; Score 17.2; DB 3; Length 1855;
Best Local Similarity 86.4%; Pred. No. 78;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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Qy 1 AGCCCTCTTCAGTCCCAATCAA 22
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Db 1502 AGCTTTTCAGTCCATTCAG 1481

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RESULT 7
US-09-536-784-71/c
; Sequence 71, Application US/09536784
; Patent No. 6573082
; GENERAL INFORMATION:
; APPLICANT: Choi et. al.
; TITLE OF INVENTION: Streptococcus pneumoniae Antigens and Vaccines
; NUMBER OF SEQUENCES: 452
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: Maryland
; COUNTRY: USA
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage
; OPERATING SYSTEM: HP Vectra 486/33
; SOFTWARE: MSDOS version 6.2
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/536,784
; FILING DATE: 30-Oct-1997
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/961,083
; FILING DATE: OCT-30-1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Michelle S. Marks
; REGISTRATION NUMBER: 41,971
; REFERENCE/DOCKET NUMBER: PB340P3
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (301) 309-8504
; TELEFAX: (301) 309-8512
; INFORMATION FOR SEQ ID NO: 71:
; SEQUENCE CHARACTERISTICS:

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;
; LENGTH: 1855 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 71:
US-09-536-784-71

Query Match          78.2%; Score 17.2; DB 4; Length 1855;
Best Local Similarity 86.4%; Pred. No. 78;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
Db 1502 AGCCTTTTCAGTCCCAATCAAG 1481

RESULT 8
US-09-583-110-1422/c
; Sequence 1422, Application US/09583110
; Patent No. 6699703
; GENERAL INFORMATION:
; APPLICANT: Lynn Doucette-Stamm et al.
; TITLE OF INVENTION: Nucleic Acid and Amino Acid Sequences Relating to Streptococcus
; FILE REFERENCE: Pneumoniae for Diagnostics and Therapeutics
; PATH00-07A
; CURRENT APPLICATION NUMBER: US/09/583,110
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 09/107,433
; PRIOR FILING DATE: 1998-06-30
; PRIOR APPLICATION NUMBER: US 60/085,131
; PRIOR FILING DATE: 1998-05-12
; PRIOR APPLICATION NUMBER: US 60/051,553
; PRIOR FILING DATE: 1997-07-02
; NUMBER OF SEQ ID NOS: 5322
; SEQ ID NO 1422
; LENGTH: 3615
; TYPE: DNA
; ORGANISM: Streptococcus pneumoniae
US-09-583-110-1422

Query Match          78.2%; Score 17.2; DB 4; Length 3615;
Best Local Similarity 86.4%; Pred. No. 89;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
Db 3181 AGCCTTTTCAGTCCCAATCAAG 3160

RESULT 9
US-09-107-433-2464/c
; Sequence 2464, Application US/09107433
; Patent No. 6800744
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID
; SEQUENCES RELATING TO STREPTOCOCCUS PNEUMONIAE
; THERAPEUTICS
; NUMBER OF SEQUENCES: 5206
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD/ROM ISO9660
; COMPUTER: <Unknown>
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: <Unknown>
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,433
; FOR DIAGNO
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; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/ 085131
; FILING DATE: May 12, 1998
; APPLICATION NUMBER: 60/051553
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Atinello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-011
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 2464:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 3789 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Streptococcus pneumoniae
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (B) LOCATION 1...3789
; SEQUENCE DESCRIPTION: SEQ ID NO: 2464:
US-09-107-433-2464

Query Match          78.2%; Score 17.2; DB 4; Length 3789;
Best Local Similarity 86.4%; Pred. No. 90;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
Db 3355 AGCCTTTTCAGTCCCAATCAAG 3334

RESULT 10
US-10-144-198-13
; Sequence 13, Application US/10144198
; Patent No. 6833247
; GENERAL INFORMATION:
; APPLICANT: Origene Technologies Inc
; TITLE OF INVENTION: Regulated Prostate Cance Genes
; FILE REFERENCE: 9U 105 R1
; CURRENT APPLICATION NUMBER: US/10/144,198
; CURRENT FILING DATE: 2002-05-14
; NUMBER OF SEQ ID NOS: 44
; SOFTWARE: Patentin version 3.0
; SEQ ID NO 13
; LENGTH: 4191
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (121)..(3246)
US-10-144-198-13

Query Match          78.2%; Score 17.2; DB 4; Length 4191;
Best Local Similarity 86.4%; Pred. No. 92;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
Db 831 ACCCCTCTTCAGCCCCATCAAG 852

RESULT 11
US-08-961-527-74/c
; Sequence 74, Application US/08961527
; Patent No. 6420135
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GENERAL INFORMATION:
APPLICANT: Charles Kunsch
TITLE OF INVENTION: Streptococcus pneumoniae Polynucleotides and Sequences
NUMBER OF SEQUENCES: 391
CORRESPONDENCE ADDRESS:
ADDRESSEE: Human Genome Sciences, Inc.
STREET: 9410 Key West Avenue
CITY: Rockville
STATE: Maryland
COUNTRY: USA
ZIP: 20850
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage
COMPUTER: HP Vectra 486/33
OPERATING SYSTEM: MSDOS version 6.2
SOFTWARE: ASCII Text
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/961,527
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Brookes, A. Anders
REGISTRATION NUMBER: 36,373
REFERENCE/DOCKET NUMBER: PB340P1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (301) 309-8504
TELEFAX: (301) 309-8512
INFORMATION FOR SEQ ID NO: 74:
SEQUENCE CHARACTERISTICS:
LENGTH: 16535 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
US-08-961-527-74

Query Match 78.2%; Score 17.2; DB 3; Length 16535;
Best Local Similarity 86.4%; Pred. No. 1.2e+02;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
DB 3310 AGCCCTCTTCAGTCCCAATCAAG 3289

RESULT 12
US-09-949-016-12747
Sequence 12747, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: CL001307
CURRENT APPLICATION NUMBER: US/09/949,016
CURRENT FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 12747
LENGTH: 55328
TYPE: DNA
ORGANISM: Human
US-09-949-016-12747

Query Match 78.2%; Score 17.2; DB 4; Length 55328;

Best Local Similarity 86.4%; Pred. No. 1.5e+02;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
DB 17089 AGCCCTCTTCAGTCCCAATCAAG 17110

RESULT 13
US-09-949-016-17146
Sequence 17146, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: CL001307
CURRENT APPLICATION NUMBER: US/09/949,016
CURRENT FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 17146
LENGTH: 55330
TYPE: DNA
ORGANISM: Human
US-09-949-016-17146

Query Match 78.2%; Score 17.2; DB 4; Length 55330;
Best Local Similarity 86.4%; Pred. No. 1.5e+02;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
DB 17089 AGCCCTCTTCAGTCCCAATCAAG 17110

RESULT 14
US-09-092-437-1/c
Sequence 1, Application US/09092437
Patent No. 6190881
GENERAL INFORMATION:
APPLICANT: Wilding, Edwina Imogen
APPLICANT: Black, Michael T.
APPLICANT: Shilling, Lisa K.
APPLICANT: Kosmatka, Anna L.
APPLICANT: Jaworski, Deborah D.
APPLICANT: Wang, Min
TITLE OF INVENTION: nrdf
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: Dechert, Price & Rhoads
STREET: 4000 Bell Atlantic Tower, 1717 Arch Stre
CITY: Philadelphia
STATE: PA
COUNTRY: USA
ZIP: 19103-2793
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: FastSeq for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/092,437
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER:

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/
/
/ FILING DATE:
/ ATTORNEY/AGENT INFORMATION:
/ NAME: Falk, Stephen T
/ REGISTRATION NUMBER: 36,795
/ REFERENCE/DOCKET NUMBER: GM10155
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: 215-994-2488
/ TELEFAX: 215-994-2222
/ TELEX:
/ INFORMATION FOR SEQ ID NO: 1:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 963 base pairs
/ TYPE: nucleic acid
/ STRANDEDNESS: double
/ TOPOLOGY: linear
/
US-09-092-437-1
Query Match 74.5%; Score 16.4; DB 3; Length 963;
Best Local Similarity 94.4%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CCTCTTCAGTCCCAATCAA 21
Db 745 CCTCTTCAGTCCCAACAA 728

RESULT 15
US-09-583-110-1197/c
/ Sequence 533, Application US/09583110
/ Patent No. 6699703
/ GENERAL INFORMATION:
/ APPLICANT: Lynn Doucette-Stamm et al.
/ TITLE OF INVENTION: Nucleic Acid and Amino Acid Sequences Relating to Streptococcus
/ FILE OF INVENTION: Pneumoniae for Diagnostics and Therapeutics
/ FILE REFERENCE: PATH00-07A
/ CURRENT APPLICATION NUMBER: US/09/583,110
/ CURRENT FILING DATE: 2000-05-26
/ PRIOR APPLICATION NUMBER: US 09/107,433
/ PRIOR FILING DATE: 1998-06-30
/ PRIOR APPLICATION NUMBER: US 60/085,131
/ PRIOR FILING DATE: 1998-05-12
/ PRIOR APPLICATION NUMBER: US 60/051,553
/ PRIOR FILING DATE: 1997-07-02
/ NUMBER OF SEQ ID NOS: 5322
/ SEQ ID NO 1197
/ LENGTH: 963
/ TYPE: DNA
/ ORGANISM: Streptococcus pneumoniae
/
US-09-583-110-1197

Query Match 74.5%; Score 16.4; DB 4; Length 963;
Best Local Similarity 94.4%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CCTCTTCAGTCCCAATCAA 21
Db 745 CCTCTTCAGTCCCAACAA 728

RESULT 16
US-09-107-433-533/c
/ Sequence 533, Application US/09107433
/ Patent No. 6800744
/ GENERAL INFORMATION:
/ APPLICANT: Lynn A Doucette-Stamm and David Bush
/ TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID
/ SEQUENCES RELATING TO STREPTOCOCCUS PNEUMONIAE FOR DIAGNOSTICS AND THERAPEUTICS
/ NUMBER OF SEQUENCES: 5206
/ CORRESPONDENCE ADDRESS:
/ ADDRESSEE: GENOME THERAPEUTICS CORPORATION
/ STREET: 100 Beaver Street
/ CITY: Waltham
/
US-09-107-433-533
```

```
/
/
/ STATE: Massachusetts
/ COUNTRY: USA
/ ZIP: 02354
/ COMPUTER READABLE FORM:
/ MEDIUM TYPE: CD-ROM ISO9660
/ COMPUTER: <Unknown>
/ OPERATING SYSTEM: <Unknown>
/ SOFTWARE: <Unknown>
/ CURRENT APPLICATION DATA:
/ APPLICATION NUMBER: US/09/107,433
/ FILING DATE: 30-Jun-1998
/ PRIOR APPLICATION DATA:
/ APPLICATION NUMBER: 60/ 085131
/ FILING DATE: May 12, 1998
/ APPLICATION NUMBER: 60/051553
/ FILING DATE: July 2, 1997
/ ATTORNEY/AGENT INFORMATION:
/ NAME: Ariniello, Pamela Deneke
/ REGISTRATION NUMBER: 40,489
/ REFERENCE/DOCKET NUMBER: GTC-011
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: (781)893-5007
/ TELEFAX: (781)893-8277
/ INFORMATION FOR SEQ ID NO: 533:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 963 base pairs
/ TYPE: nucleic acid
/ STRANDEDNESS: double
/ TOPOLOGY: circular
/ MOLECULE TYPE: DNA (genomic)
/ HYPOTHETICAL: NO
/ ANTI-SENSE: NO
/ ORIGINAL SOURCE:
/ ORGANISM: Streptococcus pneumoniae
/ FEATURE:
/ NAME/KEY: misc feature
/ LOCATION: (8) LOCATION 1...963
/ SEQUENCE DESCRIPTION: SEQ ID NO: 533:
/
US-09-107-433-533
Query Match 74.5%; Score 16.4; DB 4; Length 963;
Best Local Similarity 94.4%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CCTCTTCAGTCCCAATCAA 21
Db 745 CCTCTTCAGTCCCAACAA 728

RESULT 17
US-09-324-258-6/c
/ Sequence 6, Application US/09324258
/ Patent No. 6723517
/ GENERAL INFORMATION:
/ APPLICANT: Bamdad, Cynthia C.
/ TITLE OF INVENTION: THE USE OF SELF-ASSEMBLED MONOLAYERS TO
/ TITLE OF INVENTION: PROBE THE STRUCTURE OF A TARGET MOLECULE
/ FILE REFERENCE: M1015/7004/TJO
/ CURRENT APPLICATION NUMBER: US/09/324,258
/ CURRENT FILING DATE: 1999-06-02
/ PRIOR APPLICATION NUMBER: U.S. 60/087,766
/ PRIOR FILING DATE: 1998-06-02
/ NUMBER OF SEQ ID NOS: 20
/ SOFTWARE: fastseq for Windows Version 3.0
/ SEQ ID NO 6
/ LENGTH: 3694
/ TYPE: DNA
/ ORGANISM: Saccharomyces cerevisiae
/ FEATURE:
/ NAME/KEY: CDS
/ LOCATION: (443)...(3088)
/ OTHER INFORMATION: GAL4
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US-09-324-258-6
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Query Match 74.5%; Score 16.4; DB 4; Length 3694;
Best Local Similarity 94.4%; Pred. No. 2.2e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CCTCTTCAGTCCCAATCAA 21
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Db 974 CCTCTTCAGTCCCAATCAA 957

RESULT 18
US-08-961-527-212
; Sequence 212, Application US/08961527
; Patent No. 6420135
; GENERAL INFORMATION:
; APPLICANT: Charles Kunsch
; TITLE OF INVENTION: Streptococcus pneumoniae Polynucleotides and Sequences
; NUMBER OF SEQUENCES: 391
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: Maryland
; COUNTRY: USA
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 1.4mb storage
; COMPUTER: HP Vectra 486/33
; OPERATING SYSTEM: MSDOS version 6.2
; SOFTWARE: ASCII Text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/961,527
; FILING DATE:
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Brookes, A. Anders
; REGISTRATION NUMBER: 36,373
; REFERENCE/DOCKET NUMBER: PB340P1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (301) 309-8504
; TELEFAX: (301) 309-8512
; INFORMATION FOR SEQ ID NO: 212:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 3902 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
US-08-961-527-212

Query Match 74.5%; Score 16.4; DB 3; Length 3902;
Best Local Similarity 94.4%; Pred. No. 2.2e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CCTCTTCAGTCCCAATCAA 21
|||||
Db 449 CCTCTTCAGTCCCAATCAA 466

RESULT 19
US-09-949-016-16603
; Sequence 16603, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08

; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16603
; LENGTH: 232547
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-16603

Query Match 74.5%; Score 16.4; DB 4; Length 232547;
Best Local Similarity 94.4%; Pred. No. 4.9e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CCTCTTCAGTCCCAATCAA 21
|||||
Db 190177 CCTCTTCAGTCCCAATCAA 190194

RESULT 20
US-09-949-016-145791/c
; Sequence 145791, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 145791
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-145791

Query Match 73.6%; Score 16.2; DB 4; Length 601;
Best Local Similarity 85.7%; Pred. No. 1.9e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GCCTCTTCAGTCCCAATCAAG 22
|||
Db 391 GCATCTTCAGTCCCAATCAAG 371

RESULT 21
US-09-949-016-146059/c
; Sequence 146059, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08

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; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 146059
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-146059

Query Match          73.6%; Score 16.2; DB 4; Length 601;
Best Local Similarity 85.7%; Pred. No. 1.9e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GCCTCTTCAGTCCAATCAAG 22
Db 391 GCATCCTTCAGTCCAATCAAG 371

RESULT 22
US-09-949-016-146327/c
; Sequence 146327, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 146327
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-146327

Query Match          73.6%; Score 16.2; DB 4; Length 601;
Best Local Similarity 85.7%; Pred. No. 1.9e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GCCTCTTCAGTCCAATCAAG 22
Db 391 GCATCCTTCAGTCCAATCAAG 371

RESULT 23
US-09-949-016-195128/c
; Sequence 195128, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 195128
; LENGTH: 601
; TYPE: DNA
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; ORGANISM: Human
US-09-949-016-195128

Query Match          73.6%; Score 16.2; DB 4; Length 601;
Best Local Similarity 85.7%; Pred. No. 1.9e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCAA 21
Db 64 AGCCCACTCTAGTCCAATCAA 44

RESULT 24
US-09-533-029-9
; Sequence 9, Application US/09533029
; Patent No. 6664446
; GENERAL INFORMATION:
; APPLICANT: Heard, Jacqueline
; APPLICANT: Broun, Pierre
; APPLICANT: Riechmann, Jose-Luis
; APPLICANT: Keddie, James
; APPLICANT: Pineda, Omaira
; APPLICANT: Adam, Luc
; APPLICANT: Samaha, Raymond
; APPLICANT: Zhang, James
; APPLICANT: Yu, Guo-Liang
; APPLICANT: Ratcliffe, Oliver
; APPLICANT: Pilgrim, Marsha
; APPLICANT: Jiang, Cai-Zhong
; APPLICANT: Reuber, Lynne
; TITLE OF INVENTION: DISEASE-INDUCED POLYNUCLEOTIDES
; FILE REFERENCE: MBI-010
; CURRENT APPLICATION NUMBER: US/09/533,029
; CURRENT FILING DATE: 2000-03-22
; EARLIER APPLICATION NUMBER: 60/125,814
; EARLIER FILING DATE: 1999-03-23
; NUMBER OF SEQ ID NOS: 121
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 9
; LENGTH: 627
; TYPE: DNA
; ORGANISM: Arabidopsis thaliana
; FEATURE:
; OTHER INFORMATION: G435
US-09-533-029-9

Query Match          73.6%; Score 16.2; DB 4; Length 627;
Best Local Similarity 85.7%; Pred. No. 2e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GCCTCTTCAGTCCAATCAAG 22
Db 249 GCATCTTCAGTCCAAGCAG 269

RESULT 25
US-09-640-211A-212
; Sequence 212, Application US/09640211A
; Patent No. 6833446
; GENERAL INFORMATION:
; APPLICANT: Wood, Marion
; APPLICANT: Shenk, Michael A.
; APPLICANT: McGrath, Annette
; APPLICANT: Glenn, Matthew
; TITLE OF INVENTION: Compositions and Methods for the
; MODIFICATION OF GENE TRANSCRIPTION
; FILE REFERENCE: 11000.1021C1U
; CURRENT APPLICATION NUMBER: US/09/640,211A
; CURRENT FILING DATE: 2000-08-16
; NUMBER OF SEQ ID NOS: 2368
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 212
; LENGTH: 850
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; TYPE: DNA
; ORGANISM: Eucalyptus grandis
US-09-640-211A-212
Query Match 73.6%; Score 16.2; DB 4; Length 850;
Best Local Similarity 85.7%; Pred. No. 2.1e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 2 GCCCTCTTCAGTCCAATCAAG 22
||| ||||| ||||| |||||
Db 32 GCCCTCTTCAGTCCAATCAAG 52

Search completed: September 6, 2005, 21:59:08
Job time : 69.9062 secs

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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 16:01:23 ; Search time 198.688 Seconds
(without alignments)
655.473 Million cell updates/sec

Title: US-10-729-421-35

Perfect score: 22
Sequence: 1 agcccttcagtcacatcaag 22

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 100 summaries

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1: Geneseqn1980s:.*
2: Geneseqn1990s:.*
3: Geneseqn2000s:.*
4: Geneseqn2001as:.*
5: Geneseqn2001bs:.*
6: Geneseqn2002as:.*
7: Geneseqn2002bs:.*
8: Geneseqn2003as:.*
9: Geneseqn2003bs:.*
10: Geneseqn2003cs:.*
11: Geneseqn2003ds:.*
12: Geneseqn2004as:.*
13: Geneseqn2004bs:.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	22	100.0	24	12	ADN36704 West Nile
3	22	100.0	24	12	ADN36702 West Nile
4	22	100.0	24	12	ADN36703 West Nile
5	22	100.0	51	12	ADN36716 West Nile
6	22	100.0	51	12	ADN36714 West Nile
7	22	100.0	51	12	ADN36715 West Nile
8	22	100.0	69	12	ADN36694 West Nile
9	22	100.0	365	6	ABK51710 Partial c
10	22	100.0	366	8	ABQ76684 WNV Cwt DN
11	22	100.0	967	12	ADQ30647 West Nile
12	22	100.0	10945	13	ADR32078 Genomic D
13	22	100.0	10945	13	ADR67768 West Nile
14	22	100.0	10975	12	ADN98022 West Nile
15	22	100.0	11029	8	ABZ68481 Nucleotid
16	22	100.0	11029	10	ABV74821 West Nile
17	22	100.0	11029	12	ADN98023 West Nile
18	21	95.5	22	12	ADN36705 West Nile
19	21	95.5	49	12	ADN36717 West Nile
20	20.4	92.7	51	12	ADN36713 West Nile

21	20	90.9	24	12	ADN36706	Adn36706 West Nile
22	20	90.9	51	12	ADN36718	Adn36718 West Nile
23	19	86.4	24	12	ADN36701	Adn36701 West Nile
C 24	18.8	85.5	10962	12	ADK13681	Adk13681 West Nile
C 25	18.8	85.5	44920	12	ADQ97910	Adq97910 Human can
26	18	81.8	532	1	AAH90221	Aan90221 Malaria-g
27	17.8	80.9	247	3	AAQ09236	Aac09236 Human sec
C 28	17.8	80.9	348	3	AAA43855	Aaa43855 Human sec
C 29	17.8	80.9	425	10	ADF80038	Adf80038 Leukaemia
C 30	17.8	80.9	447	2	AAV49571	Av49571 Human lym
C 31	17.8	80.9	616	13	ADS19174	Adsl9174 Human C-t
C 32	17.8	80.9	555	2	AAV83108	Av83108 Human C-t
C 33	17.8	80.9	597	2	AAV49570	Av49570 Human lym
C 34	17.8	80.9	697	2	AAV54641	Av54641 Nucleotid
C 35	17.8	80.9	697	2	ABL41987	Ab41987 Nucleotid
C 36	17.8	80.9	701	10	ADC38673	Adc38673 Human cdn
C 37	17.8	80.9	759	6	ABK84720	Abk84720 Human cdn
C 38	17.8	80.9	759	6	ABK48103	Abk48103 Human cdn
C 39	17.8	80.9	759	10	ADD18707	Add18707 Human dis
C 40	17.8	80.9	759	13	ADR06491	Adr06491 Human AIC
C 41	17.8	80.9	759	13	ADP54852	Adp54852 Human PRO
C 42	17.8	80.9	935	12	ADQ18422	Adq18422 Human sof
C 43	17.8	80.9	1404	12	ADQ22924	Adq22924 Human sof
C 44	17.8	80.9	1520	13	ADT46681	Adt46681 Bacterial
C 45	17.8	80.9	9375	4	AAK84948	Aak84948 Human imm
C 46	17.8	80.9	10301	4	AAK84949	Aak84949 Human imm
C 47	17.8	80.9	110000	2	AAV21209	Av21209 Human imm
C 48	17.2	78.2	1420	6	ABT08161	Abt08161 NLS-A reco
C 49	17.2	78.2	1441	6	ABT08163	Abt08163 NLS-A reco
C 50	17.2	78.2	1855	2	AAV27359	Av27359 Streptoco
C 51	17.2	78.2	1855	6	ABQ84827	Abq84827 S. pneumo
C 52	17.2	78.2	1855	10	ADC45152	Adc45152 S. pneumo
C 53	17.2	78.2	3119	5	AA572586	Aa572586 DNA encod
C 54	17.2	78.2	3789	13	ADR93829	Adr93829 Novel S.
C 55	17.2	78.2	3840	10	ABX05894	Abx05894 S. pneumo
C 56	17.2	78.2	3840	12	ADM91840	Adm91840 S. pneumo
C 57	17.2	78.2	4841	4	AAK52955	Aak52955 Human pol
C 58	17.2	78.2	4880	4	AAK51971	Aak51971 Human pol
C 59	17.2	78.2	4898	10	ABZ79896	Abz79896 Human nuc
C 60	17.2	78.2	4915	13	ADQ18215	Adq18215 Human sof
C 61	17.2	78.2	4915	13	ADP26068	Adp26068 Breast ca
C 62	17.2	78.2	5037	12	ADQ22765	Adq22765 Human sof
C 63	17.2	78.2	5309	6	ABT08172	Abt08172 Recombina
C 64	17.2	78.2	16080	6	ADT28651	Adt28651 Human Sal
C 65	17.2	78.2	16535	2	AAV52207	Av52207 Streptoco
C 66	17.2	78.2	110000	10	AB556454	Ab556454 2 of
C 67	17.2	78.2	163701	13	ABD33351	Abd33351 Murine ca
C 68	17	77.3	17	6	ACN05481	Acn05481 WNV Amber
C 69	17	77.3	17	6	ACN09471	Acn09471 WNV minus
C 70	17	77.3	17	6	ACN04728	Acn04728 WNV DNA2Y
C 71	17	77.3	17	6	ACN09470	Acn09470 WNV minus
C 72	17	77.3	17	6	ACN13599	Acn13599 WNV minus
C 73	17	77.3	17	6	ACN12229	Acn12229 WNV minus
C 74	17	77.3	17	6	ACN09469	Acn09469 WNV minus
C 75	17	77.3	17	6	ACN01443	Acn01443 WNV Inoxy
C 76	17	77.3	17	6	ACN05482	Acn05482 WNV Amber
C 77	16.8	76.4	342	8	ABX99067	Abx99067 Rice endo
C 78	16.8	76.4	440	6	ABQ55371	Abq55371 Human ova
C 79	16.8	76.4	594	12	ACH75379	Ach75379 Human gen
C 80	16.8	76.4	742	2	AAV00437	Av000437 Clone H90
C 81	16.8	76.4	943	5	AA566880	Aa566880 DNA encod
C 82	16.8	76.4	4547	6	ADT28652	Adt28652 Mouse Sal
C 83	16.8	76.4	143412	11	ACN44512	Acn44512 Mouse gen
C 84	16.4	74.5	518	13	ADQ79152	Adq79152 Novel can
C 85	16.4	74.5	650	12	ADK34072	Adk34072 Yeast let
C 86	16.4	74.5	656	13	ADQ50297	Adq50297 Novel can
C 87	16.4	74.5	960	10	ABX06892	Abx06892 S. pneumo
C 88	16.4	74.5	963	3	AA246474	Aa246474 S. pneumo
C 89	16.4	74.5	963	4	AA555886	Aa555886 Streptoco
C 90	16.4	74.5	963	4	AA555543	Aa555543 Streptoco
C 91	16.4	74.5	963	8	ACA49937	Ac49937 Prokaryot
C 92	16.4	74.5	963	13	ADR91898	Adr91898 Novel S.
C 93	16.4	74.5	2643	6	ABN79826	Abn79826 Fungal 2B

C 94 16.4 74.5 2646 2 AAQ61607 Mutated G
 C 95 16.4 74.5 2646 13 AD747700 Bacterial
 C 96 16.4 74.5 2811 12 ADJ92822 Saccharom
 C 97 16.4 74.5 3189 2 AAV65242 DNA encod
 C 98 16.4 74.5 3694 6 ABK86400 Yeast GAL
 C 99 16.4 74.5 3694 12 ADN60220 S. cerevi
 C 100 16.4 74.5 3902 2 AAV52345 Streptoco

ALIGNMENTS

RESULT 1
 ADQ30665
 ID ADQ30665 standard; DNA; 22 BP.
 XX
 AC ADQ30665;
 DT 23-SEP-2004 (first entry)
 XX
 DE West Nile Virus capsid gene antisense primer WNVVA2.
 KW ss; primer; West Nile Virus; diagnosis.
 XX
 OS West Nile virus.
 XX
 PN WO2004055159-A2.
 XX
 PD 01-JUL-2004.
 XX
 PF 05-DEC-2003; 2003WO-US038750.
 XX
 PR 12-DEC-2002; 2002US-0432850P.
 PR 20-JUN-2003; 2003US-0480431P.
 XX
 PA (CHIR) CHIRON CORP.
 XX
 PI Shyamala V;
 XX
 DR WPI; 2004-488058/46.
 XX
 PT New isolated oligonucleotides for accurately diagnosing West Nile virus
 PT infection or for capturing, detecting and quantitating West Nile virus in
 PT blood samples.
 XX
 PS Claim 1; SEQ ID NO 35; 56pp; English.
 XX
 CC The invention relates to an isolated oligonucleotide not more than 60
 CC nucleotides in length comprising a nucleotide sequence (S1) of at least
 CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
 CC 20, 21 or 23 bp) given in the specification derived from the West Nile
 CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
 CC identity to the nucleotide sequence of (S1), or complements of (S1) and
 CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
 CC end and/or the 3'-end. The detectable label is a fluorescent label
 CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
 CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
 CC composition and methods are useful for accurately diagnosing West Nile
 CC virus infection or for capturing, detecting and quantitating West Nile
 CC virus in biological samples, particularly blood samples. This sequence
 CC corresponds to a PCR primer to amplify a fragment of the capsid gene of
 CC the WNV genome. The fragment is detected using the oligonucleotides of
 CC the invention.
 XX
 SQ Sequence 22 BP; 6 A; 8 C; 3 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 12; Length 22;
 Best Local Similarity 100.0%; Pred. No. 0.8;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 AGCCCTCTTCAGTCCAATCAAG 22
 |||||
 Db 1 AGCCCTCTTCAGTCCAATCAAG 22

RESULT 2
 ADN36704
 ID ADN36704 standard; DNA; 24 BP.
 XX
 AC ADN36704;
 XX
 DT 15-JUL-2004 (first entry)
 XX
 DE West Nile virus detection-related oligonucleotide probe SeqID26.
 XX
 KW hybridisation assay probe; nucleic acid detection;
 KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 KW high throughput screening; probe; ss.
 XX
 OS West Nile virus.
 XX
 PN WO2004036190-A2.
 XX
 PD 29-APR-2004.
 XX
 PF 10-OCT-2003; 2003WO-US033639.
 XX
 PR 16-OCT-2002; 2002US-0418891P.
 PR 25-NOV-2002; 2002US-0429006P.
 PR 24-FEB-2003; 2003US-0449810P.
 XX
 PA (GENP-) GEN-PROBE INC.
 XX
 PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
 XX
 DR WPI; 2004-389590/36.
 XX
 PT New hybridization assay probe comprising target-complementary sequence of
 PT bases, useful in detecting flavivirus, e.g. West Nile virus.
 XX
 PS Claim 78; SEQ ID NO 26; 135pp; English.
 XX
 CC This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of an oligonucleotide probe which is related
 CC to the invention.
 XX
 SQ Sequence 24 BP; 7 A; 8 C; 3 G; 6 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 12; Length 24;
 Best Local Similarity 100.0%; Pred. No. 0.81;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCAAG 22
 |||||
 Db 3 AGCCCTCTTCAGTCCAATCAAG 24

RESULT 3
 ADN36702
 ID ADN36702 standard; DNA; 24 BP.
 XX
 AC ADN36702;
 XX
 DT 15-JUL-2004 (first entry)
 XX

```
DE West Nile virus detection-related oligonucleotide probe SeqID24.
XX hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; probe; ss.
XX West Nile virus.
OS WO2004036190-A2.
XX
XX 29-APR-2004.
XX
XX 10-OCT-2003; 2003WO-US033639.
XX
XX 16-OCT-2002; 2002US-0418891P.
XX
XX 25-NOV-2002; 2002US-0429006P.
XX
XX 24-FEB-2003; 2003US-0449810P.
XX
XX (GENP-) GEN-PROBE INC.
XX
XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX WPI; 2004-389590/36.
XX
XX New hybridization assay probe comprising target-complementary sequence of
PT bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX Claim 78; SEQ ID NO 24; 135pp; English.
XX
XX This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental
CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of an oligonucleotide probe which is related
CC to the invention.
XX
XX Sequence 24 BP; 7 A; 8 C; 4 G; 5 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 22; DB 12; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.81;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 AGCCCTCTTCAGTCCCAATCAAG 22
Db : |||||
1 AGCCCTCTTCAGTCCCAATCAAG 22

RESULT 4
ADN36703
ID ADN36703 standard; DNA; 24 BP.
XX
XX ADN36703;
AC
XX
XX 15-JUL-2004 (first entry)
DT
XX
XX West Nile virus detection-related oligonucleotide probe SeqID25.
DE
XX hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; probe; ss.
XX
XX West Nile virus.
OS WO2004036190-A2.
XX
XX 29-APR-2004.
XX
XX 10-OCT-2003; 2003WO-US033639.
XX
XX 16-OCT-2002; 2002US-0418891P.
XX
XX 25-NOV-2002; 2002US-0429006P.
XX
XX 24-FEB-2003; 2003US-0449810P.
XX
XX (GENP-) GEN-PROBE INC.
XX
XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX WPI; 2004-389590/36.
XX
XX New hybridization assay probe comprising target-complementary sequence of
PT bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX Claim 78; SEQ ID NO 24; 135pp; English.
XX
XX This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental
CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of an oligonucleotide probe which is related
CC to the invention.
XX
XX Sequence 24 BP; 7 A; 8 C; 4 G; 5 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 22; DB 12; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.81;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 AGCCCTCTTCAGTCCCAATCAAG 22
Db : |||||
1 AGCCCTCTTCAGTCCCAATCAAG 22

RESULT 5
ADN36716
ID ADN36716 standard; DNA; 51 BP.
XX
XX ADN36716;
AC
XX
XX 15-JUL-2004 (first entry)
DT
XX
XX West Nile virus detection-related oligonucleotide probe SeqID38.
DE
XX hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; probe; ss.
XX
XX West Nile virus.
OS Enterobacteria phage T7.
XX
XX Key Location/Qualifiers
FH misc_feature 1..27
FT /tag= a
FT /note= "T7 promoter sequence"
FT misc_feature 28..51
FT /tag= b
FT /note= "WNV-complimentary sequence"
XX
XX WO2004036190-A2.
XX
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PD 29-APR-2004.
XX
XX
PF 10-OCT-2003; 2003WO-US033639.
XX
XX
PR 16-OCT-2002; 2002US-0418891P.
XX
PR 25-NOV-2002; 2002US-0429006P.
XX
PR 24-FEB-2003; 2003US-0449810P.
XX
XX
PA (GENP-) GEN-PROBE INC.
XX
XX
PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX
XX
DR WPI; 2004-389590/36.
XX
XX
PT New hybridization assay probe comprising target-complementary sequence of
PT bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX
PS Disclosure; SEQ ID NO 38; 135pp; English.
XX
XX
CC This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental
CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of an oligonucleotide probe which is related
CC to the invention.
XX
XX
SQ Sequence 51 BP; 18 A; 12 C; 8 G; 13 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 12; Length 51;
Best Local Similarity 100.0%; Pred. No. 0.9;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 AGCCCTCTTCAGTCCAATCAAG 22
Db 30 AGCCCTCTTCAGTCCAATCAAG 51

RESULT 6
ADN36714
ID ADN36714 standard; DNA; 51 BP.
XX
XX
AC ADN36714;
XX
XX
DT 15-JUL-2004 (first entry)
XX
XX
DE West Nile virus detection-related oligonucleotide probe SeqID36.
XX
XX
KW hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; probe; ss.
XX
XX
OS West Nile virus.
OS Enterobacteria phage T7.
XX
XX
FH Key Location/Qualifiers
FT misc_feature 1..27
FT /*tag= a
FT /note= "T7 promoter sequence"
FT misc_feature 28..51
FT /*tag= b
FT /note= "WNV-complimentary sequence"
XX
XX
PN WO2004036190-A2.
XX
XX
PD 29-APR-2004.
XX

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XX
XX
PF 10-OCT-2003; 2003WO-US033639.
XX
XX
PR 16-OCT-2002; 2002US-0418891P.
XX
PR 25-NOV-2002; 2002US-0429006P.
XX
PR 24-FEB-2003; 2003US-0449810P.
XX
XX
PA (GENP-) GEN-PROBE INC.
XX
XX
PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX
XX
DR WPI; 2004-389590/36.
XX
XX
PT New hybridization assay probe comprising target-complementary sequence of
PT bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX
PS Disclosure; SEQ ID NO 36; 135pp; English.
XX
XX
CC This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental
CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of an oligonucleotide probe which is related
CC to the invention.
XX
XX
SQ Sequence 51 BP; 18 A; 12 C; 9 G; 12 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 12; Length 51;
Best Local Similarity 100.0%; Pred. No. 0.9;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 AGCCCTCTTCAGTCCAATCAAG 22
Db 28 AGCCCTCTTCAGTCCAATCAAG 49

RESULT 7
ADN36715
ID ADN36715 standard; DNA; 51 BP.
XX
XX
AC ADN36715;
XX
XX
DT 15-JUL-2004 (first entry)
XX
XX
DE West Nile virus detection-related oligonucleotide probe SeqID37.
XX
XX
KW hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; probe; ss.
XX
XX
OS West Nile virus.
OS Enterobacteria phage T7.
XX
XX
FH Key Location/Qualifiers
FT misc_feature 1..27
FT /*tag= a
FT /note= "T7 promoter sequence"
FT misc_feature 28..51
FT /*tag= b
FT /note= "WNV-complimentary sequence"
XX
XX
PN WO2004036190-A2.
XX
XX
PD 29-APR-2004.
XX

```

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PF 10-OCT-2003; 2003WO-US033639.
XX
XX 16-OCT-2002; 2002US-0418891P.
PR 25-NOV-2002; 2002US-0429006P.
PR 24-FEB-2003; 2003US-0449810P.
XX
XX (GENP-) GEN-PROBE INC.
XX
XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX WPI; 2004-389590/36.
XX
XX New hybridization assay probe comprising target-complementary sequence of
PT bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX Disclosure; SEQ ID NO 37; 135pp; English.
XX
XX This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental
CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of an oligonucleotide probe which is related
CC to the invention.
XX
XX Sequence 51 BP; 17 A; 12 C; 9 G; 13 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 22; DB 12; Length 51;
Best Local Similarity 100.0%; Pred. NO. 0.9; Indels 0; Gaps 0;
Matches 22; Conservative 0; Mismatches 0;
QY 1 AGCCCTCTTCAGTCCAATCAAG 22
DB 29 AGCCCTCTTCAGTCCAATCAAG 50
RESULT 8
ADN36694
ID ADN36694 standard; DNA; 69 BP.
XX
XX ADN36694;
AC
XX 15-JUL-2004 (first entry)
DT
XX
XX West Nile virus detection-related oligonucleotide probe SeqID16.
DE
XX
XX hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; probe; ss.
XX
XX West Nile virus.
OS
XX
XX WO2004036190-A2.
PN
XX
XX 29-APR-2004.
PD
XX
XX 10-OCT-2003; 2003WO-US033639.
PF
XX 16-OCT-2002; 2002US-0418891P.
PR 25-NOV-2002; 2002US-0429006P.
PR 24-FEB-2003; 2003US-0449810P.
XX
XX (GENP-) GEN-PROBE INC.
PA
XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX WPI; 2004-389590/36.
XX
XX New hybridization assay probe comprising target-complementary sequence of
PT bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX Disclosure; SEQ ID NO 37; 135pp; English.
XX
XX This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental
CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of an oligonucleotide probe which is related
CC to the invention.
XX
XX Sequence 51 BP; 17 A; 12 C; 9 G; 13 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 22; DB 12; Length 51;
Best Local Similarity 100.0%; Pred. NO. 0.9; Indels 0; Gaps 0;
Matches 22; Conservative 0; Mismatches 0;
QY 1 AGCCCTCTTCAGTCCAATCAAG 22
DB 29 AGCCCTCTTCAGTCCAATCAAG 50
RESULT 9
ABK51710/c
ID ABK51710 standard; cDNA; 365 BP.
XX
XX ABK51710;
AC
XX 27-AUG-2002 (first entry)
DT
XX
XX Partial cDNA for west nile virus capsid protein.
DE
XX
XX Human; ss; IGE leader sequence; west nile virus capsid protein;
KW RNA secondary structure; free energy; gene therapy; cancer;
KW hyperproliferative disease; autoimmune disease; rheumatoid arthritis;
KW multiple sclerosis; Sjogren's syndrome; sarcoidosis; scleroderma;
KW insulin-dependent diabetes mellitus; autoimmune thyroiditis; psoriasis;
KW reactive arthritis; ankylosing spondylitis; polymyositis; vasculitis;
KW dermatomyositis; Crohn's disease; ulcerative colitis.
XX
XX West Nile virus.
OS
XX
XX WO200229088-A2.
PN
XX
XX 11-APR-2002.
PD
XX
XX 04-OCT-2001; 2001WO-US031451.
PF
XX 04-OCT-2000; 2000US-0237885P.
PR
XX (UYPE-) UNIV PENNSYLVANIA.
PA
XX Weiner DB, Yang J;
PI
XX
XX WPI; 2002-416682/44.
DR
XX
XX Producing recombinant protein for preparing pharmaceutical compounds to
PT treat, e.g., cancers or autoimmune disorders, comprises predicting
PT secondary structure (SS) of mRNA and modifying DNA to give mRNA with SS
PT having increased free energy.
XX
XX Example 2; Fig 1; 48pp; English.
PS
XX

```

CC The invention relates to producing (M1) a protein (I) in a recombinant
 CC expression system (II) comprising: (a) predicting the secondary structure
 CC of mRNA; (b) modifying the native heterologous DNA sequence where the
 CC mRNA transcribed from the modified DNA has a secondary structure with
 CC increased free energy; and (c) using the modified DNA in (II) for
 CC production of (I). Also included are (1) an injectable pharmaceutical
 CC composition comprising a nucleic acid molecule that includes a modified
 CC coding sequence (IV) encoding a protein operably linked to regulatory
 CC elements, where (IV) comprises a higher AT or AU content relative to the
 CC AT or AU content of the native coding sequence and further comprising a
 CC pharmaceutical carrier and (2) a recombinant viral vector comprising a
 CC nucleic acid molecule that includes (IV). The method is used for
 CC producing a protein in a recombinant expression system. Use of a nucleic
 CC acid or recombinant viral vector that have modified DNA sequences to
 CC improve protein production can be used in gene therapy and for the
 CC treatment of cancers, hyperproliferative diseases, and autoimmune
 CC diseases such as rheumatoid arthritis, multiple sclerosis, Sjogren's
 CC syndrome, sarcoidosis, insulin-dependent diabetes mellitus, autoimmune
 CC thyroiditis, reactive arthritis, ankylosing spondylitis, scleroderma,
 CC polymyositis, dermatomyositis, psoriasis, vasculitis, Crohn's disease and
 CC ulcerative colitis. The present sequence is a cDNA for West Nile virus
 CC capsid protein. Fusion constructs of modified mRNA for the capsid protein
 CC and human Igs leader sequence are used in an experiment to minimise the
 CC free energy of the capsid protein mRNA. Note: The present sequence is not
 CC shown in the specification but was created using the information in
 CC figure 1 and the sequence appearing as ABK51708

XX SQ Sequence 365 BP; 103 A; 80 C; 109 G; 73 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 6; Length 365;

Best Local Similarity 100.0%; Pred. No. 1.2;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22

DB 95 AGCCCTCTTCAGTCCCAATCAAG 74

RESULT 10

ABQ76684/c

ID ABQ76684 standard; DNA; 366 BP.

AC ABQ76684;

DT 13-MAY-2003 (first entry)

DE WNVcwt DNA fragment.

XX Capsid protein; WNVcwt; mRNA secondary structure; cancer;
 XX immunosuppressive; antirheumatic; cytostatic; antiulcer; neuroprotective;
 XX antiarthritic; antidiabetic; antithyroid; antipsoriatic; virucide; gene;
 XX antiparasitic; antiallergic; gene therapy; allergen; multiple sclerosis;
 XX protective immune response; hyperproliferative cell; ulcerative colitis;
 XX hyperproliferative disease; psoriasis; autoimmune disease; psoriasis;
 XX rheumatoid arthritis; Sjogren's syndrome; autoimmune thyroiditis;
 XX insulin dependent diabetes mellitus; Crohn's disease; ds.

OS West Nile virus.

XX Key

XX CDS Location/Qualifiers

FT 1..366

FT /tag= a

FT /product= "WNVcwt"

FT /note= "no start or stop codon given"

XX US2002123099-A1.

XX 05-SEP-2002.

XX 04-OCT-2001; 2001US-00971806.

XX 04-OCT-2000; 2000US-0237885P.

XX

PA (WEIN/) WEINER D B.

XX (YANG/) YANG J.

PI Weiner DB, Yang J;

XX WPI; 2003-066795/06.

DR P-PSDB; ABG73556.

XX Producing protein in recombinant expression system involves predicting
 PT secondary structure of RNA encoding a protein and increasing free energy
 PT for the secondary structure by modifying sequence of DNA encoding the
 PT RNA.

XX Example 2; Fig 1; 25pp; English.

XX This invention describes a novel method for producing a protein by
 CC translation of mRNA from heterologous DNA sequences. The method involves
 CC predicting the secondary structure of mRNA transcribed from a native
 CC heterologous DNA sequence, modifying the sequence where mRNA transcribed
 CC from the modified DNA sequence has a secondary structure with increased
 CC free energy compared to mRNA transcribed from native DNA and using
 CC modified heterologous DNA for protein production. The products of the
 CC invention have immunosuppressive, antirheumatic, cytostatic, antiulcer,
 CC neuroprotective, antiallergic, antidiabetic, antithyroid, antipsoriatic,
 CC virucide, antiparasitic and antiallergic activity and can be used for
 CC gene therapy. The method described is useful for producing a protein in a
 CC recombinant expression system, preferably a cell free in vitro
 CC transcription and translation system, an in vitro cell expression system,
 CC a DNA construct used in direct DNA injection, or a recombinant vector for
 CC delivery of DNA to an individual. The products of the invention are
 CC useful for eliciting broad immune responses against a target protein,
 CC i.e. proteins specifically associated with pathogens such as viruses,
 CC parasites, allergens, or the individual's own abnormal cells.
 CC Compositions containing the products of the invention confer a broad
 CC based protective immune response against hyperproliferative cells that
 CC are characteristic in hyperproliferative diseases including all forms of
 CC cancer and psoriasis. Such compositions are also useful for treating
 CC individuals suffering from autoimmune diseases including rheumatoid
 CC arthritis, multiple sclerosis, Sjogren's syndrome, insulin dependent
 CC diabetes mellitus, autoimmune thyroiditis, Crohn's disease, ulcerative
 CC colitis and psoriasis. This sequence encodes the West Nile virus wild-
 CC type capsid protein described as WNVcwt in the disclosure of the
 CC invention

XX SQ Sequence 366 BP; 103 A; 81 C; 108 G; 74 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 8; Length 366;

Best Local Similarity 100.0%; Pred. No. 1.2;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22

DB 96 AGCCCTCTTCAGTCCCAATCAAG 75

RESULT 11

ADQ30647/c

ID ADQ30647 standard; DNA; 967 BP.

XX ADQ30647;

XX 23-SEP-2004 (first entry)

DE West Nile virus internal diagnosis control sequence.

XX ss; internal control; West Nile Virus; diagnosis.

XX West Nile virus.

XX WO2004055159-A2.

XX 01-JUL-2004.

XX

PF 05-DEC-2003; 2003WO-US038750.
 XX 12-DEC-2002; 2002US-0432850P.
 PR 20-JUN-2003; 2003US-0480431P.
 XX
 PA (CHIR) CHIRON CORP.
 XX
 PI Shyamala V;
 XX
 DR WPI; 2004-488058/46.
 XX
 PT New isolated oligonucleotides for accurately diagnosing West Nile virus in
 PT infection or for capturing, detecting and quantitating West Nile virus in
 PT blood samples.
 XX
 PS Claim 27; SEQ ID NO 17; 56pp; English.
 XX
 CC The invention relates to an isolated oligonucleotide not more than 60
 CC nucleotides in length comprising a nucleotide sequence (S1) of at least
 CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
 CC 20, 21 or 23 bp) given in the specification derived from the West Nile
 CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
 CC identity to the nucleotide sequence of (S1), or complements of (S1) and
 CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
 CC end and/or the 3'-end. The detectable label is a fluorescent label
 CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
 CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
 CC composition and methods are useful for accurately diagnosing West Nile
 CC virus infection or for capturing, detecting and quantitating West Nile
 CC virus in biological samples, particularly blood samples. This sequence
 CC corresponds to an internal control sequence for the detection of WNV
 CC sequences using the oligonucleotides of the invention.
 XX
 SQ Sequence 967 BP; 273 A; 206 C; 272 G; 216 T; 0 U; 0 Other;
 Query Match 100.0%; Score 22; DB 12; Length 967;
 Best Local Similarity 100.0%; Pred. No. 1.4;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 AGCCCTCTTCAGTCCCAATCAAG 22
 DB 197 AGCCCTCTTCAGTCCCAATCAAG 176
 RESULT 12
 ADR32078/c
 ID ADR32078 standard; DNA; 10945 BP.
 AC ADR32078;
 XX
 DT 18-NOV-2004 (first entry)
 XX
 DE Genomic DNA of a West Nile virus.
 XX
 KW analysis; target; real time PCR; ds; genomic.
 XX
 OS West Nile virus.
 XX
 PN WO2004072230-A2.
 XX
 PD 26-AUG-2004.
 XX
 PF 10-FEB-2004; 2004WO-US002012.
 XX
 PR 10-FEB-2003; 2003US-00361004.
 XX
 PA (CLEA-) CLEARANT INC.
 XX
 PI Mckenney K, Gillmeister L, Marlowe K, Armistead D;
 XX
 DR WPI; 2004-625843/60.
 XX
 PT Analyzing a target nucleic acid sequence in a biological material by real

PT time PCR using nucleic acid primers that are separated by at least 750
 XX nucleic acid residues in the target sequence.
 PS Disclosure; SEQ ID NO 5; 96pp; English.
 XX
 CC The invention relates to a novel method for analysing a target nucleic
 CC acid sequence in a biological material. The method comprises adding at
 CC least two nucleic acid primers that hybridise under stringent conditions
 CC to predetermined nucleic acid sequences of the target nucleic acid
 CC sequence that are separated by at least 750 nucleic acid residues,
 CC amplifying the target nucleic acid sequence by PCR, and detecting and
 CC quantifying the target nucleic acid sequence. The methods and
 CC compositions of the present invention are useful for analysing a target
 CC nucleic acid sequence in a biological material by real time PCR using
 CC nucleic acid primers that are separated by at least 750 nucleic acid
 CC residues in the target sequence. This polynucleotide sequence represents
 CC the genomic DNA of a West Nile virus used in the target analysis method
 CC of the invention.
 XX
 SQ Sequence 10945 BP; 2999 A; 2457 C; 3143 G; 2346 T; 0 U; 0 Other;
 Query Match 100.0%; Score 22; DB 13; Length 10945;
 Best Local Similarity 100.0%; Pred. No. 1.9;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 AGCCCTCTTCAGTCCCAATCAAG 22
 DB 153 AGCCCTCTTCAGTCCCAATCAAG 132
 RESULT 13
 ADR67768/c
 ID ADR67768 standard; DNA; 10945 BP.
 AC ADR67768;
 XX
 DT 18-NOV-2004 (first entry)
 XX
 DE West Nile virus DNA detected by novel detection method.
 XX
 KW ds; detection; pathogen.
 XX
 OS West Nile virus.
 XX
 PN WO2004072231-A2.
 XX
 PD 26-AUG-2004.
 XX
 PF 10-FEB-2004; 2004WO-US002013.
 XX
 PR 10-FEB-2003; 2003US-00361002.
 XX
 PA (CLEA-) CLEARANT INC.
 XX
 PI Mckenney K, Gillmeister L, Marlowe K, Armistead D;
 XX
 DR WPI; 2004-625844/60.
 XX
 PT Determining level of potentially active biological pathogens in
 PT biological material, by adding nucleic acid primer pairs to biological
 PT material, amplifying target nucleic acid by PCR, detecting and
 PT quantifying target nucleic acid.
 XX
 PS Disclosure; SEQ ID NO 5; 111pp; English.
 XX
 CC The invention relates to a method of determining (M1) level of
 CC potentially active biological pathogens in biological material, involves
 CC adding at least two nucleic acid primer pairs to biological material,
 CC amplifying target nucleic acid sequences by PCR, and detecting and
 CC quantifying target nucleic acid sequences, where quantity of the nucleic
 CC acid sequences is proportional to number of biological pathogens in
 CC biological material. (M1) is useful for determining level of potentially
 CC active biological pathogens in a biological material such as cells,

tissues, blood or blood components, proteins, enzymes, immunoglobulins, botanicals, food, ligaments, tendons, nerves, bone, teeth, skin grafts, bone marrow, heart valves, cartilage, corneas, arteries, veins, organs, lipids, carbohydrates, collagen, chitin and its derivatives, forensic samples, mummified material, human or animal remains, stem cells, islet of Langerhans cells, cells for transplantation, red blood cells, white blood cells or platelets. The biological pathogen is chosen from bacteria, viruses, fungi and single cell parasites. The biological pathogen is chosen from Aspergillus, Candida, Histoplasma, Saccharomyces, Coccidioides, Cryptococcus, Escherichia, Bacillus, Campylobacter, Helicobacter, Listeria, Clostridium, Streptococcus, Enterococcus, Staphylococcus, Brucella, Haemophilus, Salmonella, Yersinia, Pseudomonas, Serratia, Enterobacter, Klebsiella, Proteus, Citrobacter, Corynebacterium, Propionibacterium and Coxiella. The biological pathogen is chosen from Adeno-associated virus (AAV), California encephalitis virus, Coronaviruses, Coxsackievirus-A, Coxsackievirus-B, Eastern equine encephalitis virus (EEEV), Echovirus, Hantavirus, Hepatitis A virus (HAV), Hepatitis C virus (HCV), Hepatitis delta virus (HDV), Hepatitis E virus (HEV), Hepatitis G virus (HGV), HIV, Human T-lymphotrophic virus (HTLV), Influenza virus (Flu virus), Measles virus (Rubeola), Mumps virus, Norwalk virus, Parainfluenza virus, Polio virus, Rabies virus, Respiratory Syncytial virus, Rhinovirus, Rubella virus, Saint Louis encephalitis virus, Western equine encephalitis virus (WEEV), Yellow fever virus, Adenovirus, Cytomegalovirus (CMV), Epstein-Barr virus (EBV), Hepatitis B virus (HBV), Herpes simplex virus 1, Herpes simplex virus 2, Molluscum contagiosum, Papilloma virus (HPV), Smallpox virus (Variola), Vaccinia virus, Venezuelan equine encephalitis virus (VEEV), Ebola virus, West Nile virus, Human Parvovirus B19 and Rotavirus. (M1) is useful for determining the effectiveness of a sterilization process applied to a biological material. (M1) is useful in determining whether the biological pathogen is inactive or active. (M1) enables determination of whether the particular biological pathogen is present in a biological material as shown by amplification of first target sequence and whether the biological pathogen is inactive or active. (M1) enables evaluation of the effectiveness of sterilization processes, and determination of both the original level and the residual level of potentially active biological pathogens. This sequence corresponds to a West Nile virus DNA detected by the method of the invention.

XX Sequence 10945 BP; 2999 A; 2457 C; 3143 G; 2346 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 13; Length 10945;
Best Local Similarity 100.0%; Pred. No. 1.9;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCAAG 22
|||||
DB 153 AGCCCTCTTCAGTCCAATCAAG 132

RESULT 14
ADN98022/c
ID ADN98022 standard; DNA; 10975 BP.

XX AC ADN98022;

XX DT 29-JUL-2004 (first entry)

XX DE West Nile Virus isolate 2741 complete genome sequence.

XX KW de; West Nile Virus; envelope protein; glycoprotein E; flavivirus;
XX Japanese encephalitis virus; Dengue virus; St Louis encephalitis virus.

XX OS West Nile virus.

XX PN WO2004040263-A2.

XX PD 13-MAY-2004.

XX PF 31-OCT-2003; 2003WO-US034823.

XX PR 31-OCT-2002; 2002US-0422755P.

XX PR 06-JUN-2003; 2003US-0476513P.

XX (HEAL-) HEALTH RES INC.

XX PI Wong SJ, Pei-Yong S;

XX DR WPI; 2004-400223/37.

XX DR GENBANK; AF206518.

XX PT New diagnostic kit comprising West Nile Virus (WNV) envelope protein
PT reactive with antibody against WNV and cross-reactive with antibody
PT against a flavivirus, useful in diagnosing flavivirus infection caused by
PT DENV, WNV, JEV or SLEV.

XX PS Disclosure; Fig 37; 212pp; English.

XX CC The invention relates to a diagnostic kit comprising at least one
CC isolated and purified polypeptide comprising a West Nile Virus (WNV)
CC envelope (E) protein or its immunogenic fragment having a native
CC conformation or non-denatured structure and that is reactive with
CC antibodies against WNV and cross-reactive with antibodies against a
CC flavivirus. The diagnostic kit is useful in diagnosing flavivirus
CC infection caused by DENV, WNV, JEV or SLEV. This sequence corresponds to
CC the complete nucleotide sequence of the WNV isolate 2741.

XX SQ Sequence 10975 BP; 3007 A; 2460 C; 3149 G; 2359 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 12; Length 10975;
Best Local Similarity 100.0%; Pred. No. 2;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCAAG 22
|||||
DB 177 AGCCCTCTTCAGTCCAATCAAG 156

RESULT 15
ABZ68481/c
ID ABZ68481 standard; DNA; 11029 BP.

XX AC ABZ68481;

XX DT 22-APR-2003 (first entry)

XX DE Nucleotide sequence of the genome of West Nile virus IS-98-ST1.

XX KW WNV; IS-98-ST1; Flavivirus; infection; encephalitis; gene; ss.

XX OS West nile virus.

XX FH Key Location/Qualifiers
XX CDS 97..10397
XX FT /*tag= a
XX FT /product= "polyprotein"

XX PN WO200281511-A1.

XX PD 17-OCT-2002.

XX PF 04-APR-2002; 2002WO-FR001168.

XX PR 04-APR-2001; 2001FR-00004599.

XX PR 06-SEP-2001; 2001FR-00011525.

XX PA (INSP) INST PASTEUR.

XX PA (KIMR-) KIMRON VETERINARY INST.

XX PI Despres P, Deubel V, Guenet J, Drouet M, Malkinson M, Banet C;
PI Frenkiel M, Courageot M, Coulibaly F, Catteau A, Flamand M, Weber P;
PI Ceccaldi P;

XX DR WPI; 2003-058498/05.

XX DR P-PSDB; ABP70647.

PT New neurovirulent strain of West Nile virus, useful in diagnosis and
 PT screening for antiviral agents, also related nucleic acids, proteins and
 PT antibodies.

XX Claim 1; Page 34-49; 68pp; French.

XX The present sequence represents the genome of a strain of West Nile virus
 CC (WNV), designated IS-98-ST1. This strain is a neuroinvasive and
 CC neurovirulent strain of WNV. Polynucleotides and polypeptides derived
 CC from the IS-98-ST1 genome are useful for diagnosis and prognosis of
 CC Flavivirus infection, specifically WNV-mediated encephalitis. They are
 CC also useful to raise specific antibodies, for recombinant expression of
 CC WNV proteins or peptides (for diagnosis, production of antibodies and
 CC identification of specific binding partners in cells), for identifying
 CC cellular genes implicated in resistance to viral infection, and for
 CC screening for anti-Flavivirus agents

XX Sequence 11029 BP; 3019 A; 2471 C; 3167 G; 2372 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 8; Length 11029;
 Best Local Similarity 100.0%; Pred. No. 2;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
 |||||
 DB 195 AGCCCTCTTCAGTCCCAATCAAG 174

RESULT 16

ABV74821/c
 ID ABV74821 standard; DNA; 11029 BP.

XX ABV74821;

XX 28-MAR-2003 (first entry)

XX West Nile virus strain NY99-flamingo 382-99 complete genome.

XX Virucide: hepatotropic; antiinflammatory; antiviral; OAS;
 KW 2'-5'-oligoadenylate synthase; Flavivirus infection; gene; ss.

XX West Nile Virus.

XX Key Location/Qualifiers
 XX CDS 97..10398

FT /*tag= a
 FT /product= "West Nile Virus protein"

XX W0200281741-A2.

XX 17-OCT-2002.

XX 04-APR-2002; 2002WO-FR001169.

XX 04-APR-2001; 2001FR-00004598.

XX (INSP) INST PASTEUR.

XX (CNRS) CNRS CENT NAT RECH SCI.

XX Guenet J, Mashimo T, Simon-Chazottes D, Montagutelli X;

XX Frenkiel M, Despres P, Deubel V, Bonhomme F, Lucas M;

XX WPI; 2003-058566/05.

XX P-PSDB; AB998821.

XX Identifying stimulators of oligoadenylate synthase family genes, useful

XX as antiviral agents against Flavivirus, also mutated genes responsible

XX for sensitivity to virus.

XX Example 1; Page 52-67; 93pp; French.

XX The present invention relates to a method for identifying compounds (I)

XX that can stimulate a gene of the OAS (2'-5'-oligoadenylate synthase)

CC family. The method comprises: (a) inducing expression of the OAS gene in
 CC a culture of cells from a non-human mammal (flvr/flvr or flvr/flvs;
 CC indicating resistance or sensitivity to Flavivirus infection); (b)
 CC treating cells with test compound; and (c) measuring activity of OAS gene
 CC relative to a control. (I) are potentially useful as antiviral agents for
 CC treating infections by Flaviviruses (e.g. hepatitis C; dengue; yellow
 CC fever and various forms of encephalitis). Genomic OAS DNA and derived
 CC cDNA, also the encoded proteins, are useful: (a) for treating Flavivirus
 CC infection; (b) in screening for anti-flavivirus agents; and (c) for
 CC evaluating sensitivity of subjects to Flavivirus infection and their
 CC likely response to interferon treatment, e.g. to identify patients at
 CC risk of developing severe forms of such infections. The present sequence
 CC is West Nile Virus strain NY99-flamingo 382-99 (IS-98-ST1) complete
 CC genome, which was used in an example from the invention. West Nile Virus
 CC is one such Flavivirus

XX Sequence 11029 BP; 3019 A; 2471 C; 3167 G; 2372 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 10; Length 11029;
 Best Local Similarity 100.0%; Pred. No. 2;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
 |||||
 DB 195 AGCCCTCTTCAGTCCCAATCAAG 174

RESULT 17

ADN98023/c

ID ADN98023 standard; DNA; 11029 BP.

XX ADN98023;

XX 29-JUL-2004 (first entry)

XX West Nile Virus isolate 3356 complete genome sequence.

XX ds; West Nile Virus; envelope protein; glycoprotein E; flavivirus;

KW Japanese encephalitis virus; Dengue virus; St Louis encephalitis virus.

XX West Nile virus.

XX W02004040263-A2.

XX 13-MAY-2004.

XX 31-OCT-2003; 2003WO-US034823.

XX 31-OCT-2002; 2002US-0422755P.

XX 06-JUN-2003; 2003US-0476513P.

XX (HEAL-) HEALTH RES INC.

XX Wong SJ, Pei-Yong S;

XX WPI; 2004-400223/37.

XX GENBANK; AF040756.

XX New diagnostic kit comprising West Nile Virus (WNV) envelope protein
 PT reactive with antibody against WNV and cross-reactive with antibody
 PT against a flavivirus, useful in diagnosing flavivirus infection caused by

XX DENV, WNV, JEV or SLEV.

XX Disclosure; Fig 38; 212pp; English.

XX The invention relates to a diagnostic kit comprising at least one

CC isolated and purified polypeptide comprising a West Nile Virus (WNV)

CC envelope (E) protein or its immunogenic fragment having a native

CC conformation or non-denatured structure and that is reactive with

CC antibodies against WNV and cross-reactive with antibodies against a

CC flavivirus. The diagnostic kit is useful in diagnosing flavivirus

CC infection caused by DENV, WNV, JEV or SLEV. This sequence corresponds to

CC the complete nucleotide sequence of the WNV isolate 3356.

```
XX SQ Sequence 11029 BP; 3017 A; 2466 C; 3172 G; 2374 T; 0 U; 0 Other;
Query Match 100.0%; Score 22; DB 12; Length 11029;
Best Local Similarity 100.0%; Pred. No. 2;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCAAG 22
Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 18
ADN36705
ID ADN36705 standard; DNA; 22 BP.
XX AC ADN36705;
XX DT 15-JUL-2004 (first entry)
XX DE West Nile virus detection-related PCR primer SeqID27.
XX KW hybridisation assay probe; nucleic acid detection;
XX KW target-complementary sequence; flavivirus; West Nile virus; WNV;
XX KW RNA virus; infection; meningitis; encephalitis;
XX KW high throughput screening; PCR; primer; ss.
XX OS West Nile virus.
XX PN WO2004036190-A2.
XX PD 29-APR-2004.
XX PF 10-OCT-2003; 2003WO-US033639.
XX PR 16-OCT-2002; 2002US-0418891P.
XX PR 25-NOV-2002; 2002US-0429006P.
XX PR 24-FEB-2003; 2003US-0449810P.
XX PA (GENP-) GEN-PROBE INC.
XX PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX DR WPI; 2004-389590/36.
XX PT New hybridization assay probe comprising target-complementary sequence of
XX PT bases, useful in detecting flavivirus, e.g. West Nile virus.
XX PS Example 2; SEQ ID NO 27; 135pp; English.
XX CC This invention relates to a novel hybridisation assay probe, for
XX CC detecting a nucleic acid, which is a probe sequence that comprises a
XX CC target-complementary sequence of bases, and optionally one or more base
XX CC sequences that are not complementary to the nucleic acid that is to be
XX CC detected. The hybridisation assay probes and the kits are useful in
XX CC detecting and amplifying a target nucleic acid sequence, for example
XX CC flavivirus like West Nile virus, that may be present in a biological
XX CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX CC birds and culex mosquitoes, with humans and horses serving as incidental
XX CC hosts. Infection of humans can lead to meningitis or encephalitis. The
XX CC invention may allow for accurate and efficient high throughput screening.
XX CC The present sequence is that of a PCR primer which is related to the
XX CC invention.
XX SQ Sequence 22 BP; 6 A; 8 C; 2 G; 6 T; 0 U; 0 Other;
Query Match 95.5%; Score 21; DB 12; Length 22;
Best Local Similarity 100.0%; Pred. No. 2.4;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCAAA 21
Db 2 AGCCCTCTTCAGTCCAATCAAA 22

us-10-729-421-35.rng
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```
XX SQ Sequence 11029 BP; 3017 A; 2466 C; 3172 G; 2374 T; 0 U; 0 Other;
Query Match 100.0%; Score 22; DB 12; Length 11029;
Best Local Similarity 100.0%; Pred. No. 2;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCAAG 22
Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 19
ADN36717
ID ADN36717 standard; DNA; 49 BP.
XX AC ADN36717;
XX DT 15-JUL-2004 (first entry)
XX DE West Nile virus detection-related oligonucleotide probe SeqID39.
XX KW hybridisation assay probe; nucleic acid detection;
XX KW target-complementary sequence; flavivirus; West Nile virus; WNV;
XX KW RNA virus; infection; meningitis; encephalitis;
XX KW high throughput screening; probe; ss.
XX OS West Nile virus.
XX PN WO2004036190-A2.
XX PD 29-APR-2004.
XX PF 10-OCT-2003; 2003WO-US033639.
XX PR 16-OCT-2002; 2002US-0418891P.
XX PR 25-NOV-2002; 2002US-0429006P.
XX PR 24-FEB-2003; 2003US-0449810P.
XX PA (GENP-) GEN-PROBE INC.
XX PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX DR WPI; 2004-389590/36.
XX PT New hybridization assay probe comprising target-complementary sequence of
XX PT bases, useful in detecting flavivirus, e.g. West Nile virus.
XX PS Disclosure; SEQ ID NO 39; 135pp; English.
XX CC This invention relates to a novel hybridisation assay probe, for
XX CC detecting a nucleic acid, which is a probe sequence that comprises a
XX CC target-complementary sequence of bases, and optionally one or more base
XX CC sequences that are not complementary to the nucleic acid that is to be
XX CC detected. The hybridisation assay probes and the kits are useful in
XX CC detecting and amplifying a target nucleic acid sequence, for example
XX CC flavivirus like West Nile virus, that may be present in a biological
XX CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX CC birds and culex mosquitoes, with humans and horses serving as incidental
XX CC hosts. Infection of humans can lead to meningitis or encephalitis. The
XX CC invention may allow for accurate and efficient high throughput screening.
XX CC The present sequence is that of an oligonucleotide probe which is related
XX CC to the invention.
XX SQ Sequence 49 BP; 17 A; 12 C; 7 G; 13 T; 0 U; 0 Other;
Query Match 95.5%; Score 21; DB 12; Length 49;
Best Local Similarity 100.0%; Pred. No. 2.7;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCAA 21
Db 29 AGCCCTCTTCAGTCCAATCAA 49
```

```

RESULT 20
ADN36713
ID ADN36713 standard; DNA; 51 BP.
XX
AC ADN36713;
XX
DT 15-JUL-2004 (first entry)
XX
DE West Nile virus detection-related PCR primer SeqID35.
XX
KW hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; PCR; primer; ss.
XX
OS West Nile virus.
XX
OS Enterobacteria phage T7.
XX
FH Key Location/Qualifiers
FT misc_feature 1..27
FT /*tag= a
FT /note= "T7 promoter sequence"
FT misc_feature 28..51
FT /*tag= b
FT /note= "WNV-complementary sequence"
XX
PN WO2004036190-A2.
XX
PD 29-APR-2004.
XX
PF 10-OCT-2003; 2003WO-US033639.
XX
PR 16-OCT-2002; 2002US-0418891P.
XX
PR 25-NOV-2002; 2002US-0429006P.
XX
PR 24-FEB-2003; 2003US-0449810P.
XX
PA (GENP-) GEN-PROBE INC.
XX
PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX WPI; 2004-389590/36.
XX
PT New hybridization assay probe comprising target-complementary sequence of
PT bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
PS Example 5; SEQ ID NO 35; 135pp; English.
XX
CC This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental
CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of a PCR primer which is related to the
CC invention.
XX
SQ Sequence 51 BP; 19 A; 12 C; 8 G; 12 T; 0 U; 0 Other;
Query Match 92.7%; Score 20.4; DB 12; Length 51;
Best Local Similarity 95.5%; Pred. No. 5.3;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCAAG 22
DB 25 AGACCTCTTCAGTCCAATCAAG 46

RESULT 21
ADN36706
ID ADN36706 standard; DNA; 24 BP.
XX
AC ADN36706;
XX
DT 15-JUL-2004 (first entry)
XX
DE West Nile virus detection-related PCR primer SeqID28.
XX
KW hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; PCR; primer; ss.
XX
OS West Nile virus.
XX
PN WO2004036190-A2.
XX
PD 29-APR-2004.
XX
PF 10-OCT-2003; 2003WO-US033639.
XX
PR 16-OCT-2002; 2002US-0418891P.
XX
PR 25-NOV-2002; 2002US-0429006P.
XX
PR 24-FEB-2003; 2003US-0449810P.
XX
PA (GENP-) GEN-PROBE INC.
XX
PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX WPI; 2004-389590/36.
XX
PT New hybridization assay probe comprising target-complementary sequence of
PT bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
PS Claim 78; SEQ ID NO 28; 135pp; English.
XX
CC This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental
CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of a PCR primer which is related to the
CC invention.
XX
SQ Sequence 24 BP; 7 A; 9 C; 2 G; 6 T; 0 U; 0 Other;
Query Match 90.9%; Score 20; DB 12; Length 24;
Best Local Similarity 100.0%; Pred. No. 7.5;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCA 20
DB 5 AGCCCTCTTCAGTCCAATCA 24

RESULT 22
ADN36718
ID ADN36718 standard; DNA; 51 BP.
XX
AC ADN36718;
XX
DT 15-JUL-2004 (first entry)
XX
DE West Nile virus detection-related PCR primer SeqID40.
XX

```

KW hybridisation assay probe; nucleic acid detection;
 KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 KW high throughput screening; PCR; primer; ss.

OS West Nile virus.
 OS Enterobacteria phage T7.

XX Key Location/Qualifiers
 FH misc_feature 1..27 /*tag= a
 FT /*note= "T7 promoter sequence"
 FT 28..51
 FT /*tag= b
 FT /*note= "WNV-complimentary sequence"

XX WO2004036190-A2.

PN 29-APR-2004.

XX 10-OCT-2003; 2003WO-US033639.

XX 16-OCT-2002; 2002US-0418891P.

XX 25-NOV-2002; 2002US-0429006P.

XX 24-FEB-2003; 2003US-0449810P.

XX (GENP-) GEN-PROBE INC.

XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;

XX WPI; 2004-389590/36.

XX New hybridization assay probe comprising target-complementary sequence of

XX bases, useful in detecting flavivirus, e.g. West Nile virus.

XX Example 6; SEQ ID NO 40; 135pp; English.

XX This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of a PCR primer which is related to the
 CC invention.

XX Sequence 51 BP; 18 A; 13 C; 7 G; 13 T; 0 U; 0 Other;

Query Match 90.9%; Score 20; DB 12; Length 51;
 Best Local Similarity 100.0%; Pred. No. 8.3;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCA 20
 |||||
 Db 32 AGCCCTCTTCAGTCCCAATCA 51

RESULT 23

ADN36701

ID ADN36701 standard; DNA; 24 BP.

XX ADN36701;

XX 15-JUL-2004 (first entry)

XX West Nile virus detection-related PCR primer SeqID23.

XX hybridisation assay probe; nucleic acid detection;

KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 KW high throughput screening; PCR; primer; ss.

OS West Nile virus.

PN WO2004036190-A2.

XX 29-APR-2004.

XX 10-OCT-2003; 2003WO-US033639.

XX 16-OCT-2002; 2002US-0418891P.

XX 25-NOV-2002; 2002US-0429006P.

XX 24-FEB-2003; 2003US-0449810P.

XX (GENP-) GEN-PROBE INC.

XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;

XX WPI; 2004-389590/36.

XX New hybridization assay probe comprising target-complementary sequence of

XX bases, useful in detecting flavivirus, e.g. West Nile virus.

XX Example 2; SEQ ID NO 23; 135pp; English.

XX This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of a PCR primer which is related to the
 CC invention.

XX Sequence 24 BP; 8 A; 8 C; 3 G; 5 T; 0 U; 0 Other;

Query Match 86.4%; Score 19; DB 12; Length 24;
 Best Local Similarity 100.0%; Pred. No. 23;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 CCTCTTCAGTCCCAATCAAG 22
 |||||
 Db 1 CCTCTTCAGTCCCAATCAAG 19

RESULT 24

ADK13681/c

ID ADK13681 standard; DNA; 10962 BP.

XX ADK13681;

XX 20-MAY-2004 (first entry)

XX West Nile Virus DNA sequence, SEQ ID 1.

XX Virucide; Immunostimulant; flavivirus;

KW envelope protein domain III polypeptide; envelope protein; gene; ss.

XX West Nile virus.

XX Key Location/Qualifiers

FT CDS 97..10389

FT /*tag= a

FT /*product= "West Nile Virus protein"

XX WO2004016586-A2.

XX PD 26-FEB-2004.
XX PF 18-AUG-2003; 2003WO-US025681.
XX PR 16-AUG-2002; 2002US-0403893P.
XX PR 06-FEB-2003; 2003US-0445581P.
XX PA (TEXA) UNIV TEXAS SYSTEM.
XX PI Barrett A, Beasley D, Holbrook M;
XX WPI; 2004-203756/19.
XX DR P-PSDB; ADK13682.
XX PR Diagnosing flavivirus infection by contacting a sample from a human or
PT animal with a flavivirus envelope protein domain III polypeptide, and
PT detecting formation of an immunocomplex between the envelope protein and
PT antibodies in the sample.
XX PS Disclosure; SEQ ID NO 1; 110pp; English.
XX PS The present invention relates to a method for screening for a flavivirus
CC in a subject or animal host. The method comprises: contacting a sample
CC from the subject with a composition comprising a flavivirus envelope
CC protein domain III polypeptide (ADK13683-ADK13701) under conditions that
CC permit formation of specific immunocomplex between an antibody in the
CC sample and the envelope protein domain III polypeptide; and detecting
CC whether a specific immunocomplex is formed. The present sequence is the
CC coding sequence for West Nile Virus protein, from which E protein
CC envelope protein domain III polypeptide (ADK13683) is derived.
XX SQ Sequence 10962 BP; 2997 A; 2497 C; 3100 G; 2368 T; 0 U; 0 Other;
Query Match 85.5%; Score 18.8; DB 12; Length 10962;
Best Local Similarity 90.9%; Pred. No. 68;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
DB 195 AGCCCTCTTATGCTCTATCAAG 174
RESULT 25
ADQ97910/c
ID ADQ97910 standard; DNA; 44920 BP.
XX AC ADQ97910;
XX DT 07-OCT-2004 (first entry)
XX DE Human cancer associated sequence HD11-022, SEQ ID 887.
XX KW Cytostatic; Gene Therapy; cancer; leukemia; lymphoma; Human; ds.
XX OS Homo sapiens.
XX PN WO2004060304-A2.
XX PD 22-JUL-2004.
XX PF 22-DEC-2003; 2003WO-US041389.
XX PR 27-DEC-2002; 2002US-00330773.
XX PA (SAGR-) SAGRES DISCOVERY INC.
XX PI Morris DW, Malandro MS;
XX WPI; 2004-543781/52.
XX New isolated cancer associated nucleic acids comprising at least 10
PT contiguous nucleotides, useful for diagnosing, preventing and/or treating

PT cancers such as leukemia and lymphoma.
XX Claim 1; SEQ ID NO 887; 199pp; English.
XX CC The present invention relates to cancer associated sequences (ADQ97025-
CC ADQ98004). The sequences are useful for the diagnosis, prevention and/or
CC treatment of cancer, such as leukemia and lymphoma. Note: The sequence
CC data for this patent did not form part of the printed specification, but
CC was obtained in electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences.
XX SQ Sequence 44920 BP; 13235 A; 9200 C; 9459 G; 13026 T; 0 U; 0 Other;
Query Match 85.5%; Score 18.8; DB 12; Length 44920;
Best Local Similarity 90.9%; Pred. No. 84;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
DB 42318 AGCCCTCTTCAATCTAATCAAG 42297
Search completed: September 6, 2005, 20:39:28
Job time : 202.688 secs

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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 16:01:23 ; Search time 770.688 Seconds
(without alignments)
1383.200 Million cell updates/sec

Title: US-10-729-421-35

Perfect score: 22

Sequence: 1 agccctctcagtcacatcaag 22

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 100 summaries

Database :

GenEmbl.*

1: gb_ba.*

2: gb_hgt.*

3: gb_in.*

4: gb_om.*

5: gb_ov.*

6: gb_pat.*

7: gb_ph.*

8: gb_pl.*

9: gb_pr.*

10: gb_ro.*

11: gb_sts.*

12: gb_sy.*

13: gb_un.*

14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	22	100.0	1648	14	AF375042 West Nile
C 2	22	100.0	1648	14	AF375044 West Nile
C 3	22	100.0	1648	14	AF375223 West Nile
C 4	22	100.0	2440	14	AF194117 West Nile
C 5	22	100.0	10945	14	AF202541 West Nile
C 6	22	100.0	10975	14	AF206518 West Nile
C 7	22	100.0	10989	14	AY268133 West Nile
C 8	22	100.0	10998	14	AY278441 West Nile
C 9	22	100.0	11029	6	AX576542 Sequence
C 10	22	100.0	11029	6	AX577796 Sequence
C 11	22	100.0	11029	14	AB185914 West Nile
C 12	22	100.0	11029	14	AB185915 West Nile
C 13	22	100.0	11029	14	AB185916 West Nile
C 14	22	100.0	11029	14	AB185917 West Nile
C 15	22	100.0	11029	14	AF196835 West Nile
C 16	22	100.0	11029	14	AF260967 West Nile
C 17	22	100.0	11029	14	AF404753 West Nile
C 18	22	100.0	11029	14	AF404754 West Nile
C 19	22	100.0	11029	14	AF404755 West Nile

C 20	22	100.0	11029	14	AF404756
C 21	22	100.0	11029	14	AF481864
C 22	22	100.0	11029	14	AF533540
C 23	22	100.0	11029	14	AF533540
C 24	22	100.0	11029	14	AY289214
C 25	21	95.5	1648	14	AY688948
C 26	21	95.5	1648	14	AY688948
C 27	21	95.5	2323	14	AF375043
C 28	21	95.5	2323	14	AF375043
C 29	21	95.5	10842	14	AF130362
C 30	21	95.5	10845	14	AF130363
C 31	21	95.5	10972	14	AF278442
C 32	21	95.5	10984	14	AF277252
C 33	21	95.5	10989	14	AF317203
C 34	21	95.5	11028	14	AY262283
C 35	21	95.5	11029	14	AY268132
C 36	21	95.5	11029	14	AY490240
C 37	21	95.5	11029	14	AF260968
C 38	19.4	88.2	10664	14	AF260969
C 39	19.4	88.2	11022	14	AF404757
C 40	19.4	88.2	11022	14	KUNCG
C 41	19.4	88.2	115637	9	AY274504
C 42	19.4	88.2	120769	9	AY274505
C 43	18.8	85.5	240	14	HS020
C 44	18.8	85.5	10741	14	AL662864
C 45	18.8	85.5	10962	14	WNP42SAA
C 46	18.8	85.5	133888	2	AY277251
C 47	18.8	85.5	142560	2	AC010769
C 48	18.8	85.5	152037	9	AC021906
C 49	18.8	85.5	176827	9	HS167A19
C 50	18.8	85.5	211550	9	AC022306
C 51	18.4	83.6	203841	2	AC073912
C 52	18.4	83.6	221153	2	AC141954
C 53	18.4	83.6	238643	2	AC115962
C 54	18.4	83.6	275209	2	AC133255
C 55	18.8	81.8	532	6	AC097164
C 56	17.8	80.9	247	6	E64859
C 57	17.8	80.9	247	6	AX897448
C 58	17.8	80.9	380	6	BD032981
C 59	17.8	80.9	380	6	AR424247
C 60	17.8	80.9	380	6	AX984941
C 61	17.8	80.9	380	6	BD119800
C 62	17.8	80.9	447	6	AX778437
C 63	17.8	80.9	487	6	BD063596
C 64	17.8	80.9	547	9	CQ687751
C 65	17.8	80.9	551	9	AY142147
C 66	17.8	80.9	597	6	BC033359
C 67	17.8	80.9	697	6	BD141462
C 68	17.8	80.9	697	6	E17140
C 69	17.8	80.9	697	9	BD063621
C 70	17.8	80.9	699	11	BV180382
C 71	17.8	80.9	717	9	BC005254
C 72	17.8	80.9	756	6	CQ727154
C 73	17.8	80.9	759	6	AX394284
C 74	17.8	80.9	759	9	HSALICL
C 75	17.8	80.9	820	3	AF036902
C 76	17.8	80.9	1079	3	AF061750
C 77	17.8	80.9	15284	1	U67462
C 78	17.8	80.9	16212	9	HSJ593K13
C 79	17.8	80.9	74625	9	AL596214
C 80	17.8	80.9	101016	9	AL355294
C 81	17.8	80.9	110000	6	AR271569
C 82	17.8	80.9	110000	8	CR382127
C 83	17.8	80.9	110185	9	AC079795
C 84	17.8	80.9	114428	9	AL844525
C 85	17.8	80.9	116061	9	AL732324
C 86	17.8	80.9	119430	8	ATF9D16
C 87	17.8	80.9	131552	9	AL137063
C 88	17.8	80.9	138232	2	AC012582
C 89	17.8	80.9	145820	9	AC020655
C 90	17.8	80.9	147327	9	AC097455
C 91	17.8	80.9	158560	9	AC098969
C 92	17.8	80.9	161691	2	AC021385

93 17.8 80.9 165067 9 AL138927 Human DNA
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c 95 17.8 80.9 169483 9 AC093829
c 96 17.8 80.9 171462 9 AC117832 Homo sapi
c 97 17.8 80.9 171518 2 AL589697
c 98 17.8 80.9 171710 2 AC013388 Homo sapi
c 99 17.8 80.9 173031 9 AL359853 Human DNA
100 17.8 80.9 173169 9 AC068538 Homo sapi

ALIGNMENTS

RESULT 1
AF375042/c
LOCUS
DEFINITION West Nile virus isolate WN_0043 polyprotein mRNA, partial cds.
ACCESSION AF375042
VERSION AF375042.1 GI:19421847
KEYWORDS
SOURCE West Nile virus
ORGANISM West Nile virus
Viruses; sRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
REFERENCE 1 (bases 1 to 1648)
AUTHORS Hindiyyeh,M., Shulman,L.M., Mendelson,E., Weiss,L., Grossman,Z. and Bin,H.
TITLE Isolation and characterization of West Nile virus from the blood of viremic patients during the 2000 outbreak in Israel
JOURNAL Emerging Infect. Dis. 7 (4), 748-750 (2001)
MEDLINE 21469825
PUBMED 11585544
REFERENCE 2 (bases 1 to 1648)
AUTHORS Hindiyyeh,M., Shulman,L.M., Mendelson,E., Grossman,Z., Weiss,L. and Bin,H.
TITLE Direct Submission
JOURNAL Submitted (30-APR-2001) Central Virology Laboratory, Ministry of Health, Public Health Laboratories, Sheba Medical Center, Tel Hashomer 52621, Israel
FEATURES
source Location/Qualifiers
1. .1648
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CDS
1 AGCCCTCTTCAGTCCCAATCAAG 22
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26 AGCCCTCTTCAGTCCCAATCAAG 5

ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 1648;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGCCCTCTTCAGTCCCAATCAAG 22
|||||
Db 26 AGCCCTCTTCAGTCCCAATCAAG 5
RESULT 2

AF375044/c
LOCUS
DEFINITION West Nile virus isolate WN_0247 polyprotein mRNA, partial cds.
ACCESSION AF375044
VERSION AF375044.1 GI:19421851
KEYWORDS
SOURCE West Nile virus
ORGANISM West Nile virus
Viruses; sRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
REFERENCE 1 (bases 1 to 1648)
AUTHORS Hindiyyeh,M., Shulman,L.M., Mendelson,E., Weiss,L., Grossman,Z. and Bin,H.
TITLE Isolation and characterization of West Nile virus from the blood of viremic patients during the 2000 outbreak in Israel
JOURNAL Emerging Infect. Dis. 7 (4), 748-750 (2001)
MEDLINE 21469825
PUBMED 11585544
REFERENCE 2 (bases 1 to 1648)
AUTHORS Hindiyyeh,M., Shulman,L.M., Mendelson,E., Grossman,Z., Weiss,L. and Bin,H.
TITLE Direct Submission
JOURNAL Submitted (30-APR-2001) Central Virology Laboratory, Ministry of Health, Public Health Laboratories, Sheba Medical Center, Tel Hashomer 52621, Israel
FEATURES
source Location/Qualifiers
1. .1648
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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 1648;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGCCCTCTTCAGTCCCAATCAAG 22
|||||
Db 26 AGCCCTCTTCAGTCCCAATCAAG 5
RESULT 3
AF375223/c
LOCUS
DEFINITION West Nile virus polyprotein gene, partial cds.
ACCESSION AF375223
VERSION AF375223.1 GI:17226060
KEYWORDS
SOURCE West Nile virus
ORGANISM West Nile virus
Viruses; sRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
REFERENCE 1 (bases 1 to 1648)
AUTHORS Banet,C., Brill,A., Samina,I., Yadin,H., Straum,Y., Weissman,J., Pokamonski,S., King,R., Deubel,V. and Malkinson,M.
TITLE Phylogenetic relationships of West Nile viruses isolated in Israel

JOURNAL REFERENCE AUTHORS	from 1997 to 2000 Unpublished 2 (bases 1 to 1648) Banet,C., Brill,A., Samina,I., Yadin,H., Straum,Y., Weisman,J., Pokamonski,S., King,R., Deubel,V. and Malkinson,M.
TITLE JOURNAL	Direct Submission Submitted (01-MAY-2001) Kimron Veterinary Institute, Beit Degan 50250, Israel
FEATURES source	Location/Qualifiers 1..1648 /organism="West Nile virus" /viroion /mol_type="genomic RNA" /specific_host="white eyed gull" /db_xref="taxon:11082" <1..->1648 /codon_start=3 /product="polyprotein" /protein_id="AAL37596.1" /db_xref="GI:17226061" /translation="SLIGLKRAMLSLIDGKPIRFLVALLAFFRTAIAPTRAVLDWR RGWQTKAMHLLSPFKELGTLSAINRRSKOKRGKGKTGIAVMIGLIASVGAVTIS NFQCKVTNATVDVTIPTAGKNLCIVRAMDGYMCDDTITVECPVLSDNDP EIDWCNKTSAYYGRYCGRTKRHSRRSRSLTVQHGESLANKKGANWDSTKATRY LVKTSWLNRNGYALVAAVICGMGLSNMTGVFVLLLPAIPAYSFCNLGMSNRDF LEGVSATWDLVLDEGSCVTIMSKOKEPTIDVKMMMEANLAERYSICYLATVSDLS TKAACPTGEAHNRADPAFVRCQGVDRGNMGCGILFGKSIDTCAKFACTKAIG RTILKENIKYEVAIFVHGPTTVESHGNYSTOVGATOAGRFSITPAAPSYYTLKLGEYE VVDCPEPSGIDTIYVMTVGTFTFLVHREWFMDLNLPSSAGSTVVRNRETLMEE EPHATKQSIALGSOGEGHALAGAIPIVFESSNTVKLTSG"
CDS	Query Match 100.0%; Score 22; DB 14; Length 1648; Best Local Similarity 100.0%; Pred. No. 1.5; Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
ORIGIN	Qy 1 AGCCCTCTTCAGTCCAATCAAG 22 Db 26 AGCCCTCTTCAGTCCAATCAAG 5
LOCUS	AF194117 2440 bp RNA linear VRL 19-JAN-2000
DEFINITION	West Nile virus structural protein precursor, gene, partial cds.
ACCESSION	AF194117
VERSION	AF194117.1 GI:6715269
SOURCE	West Nile virus
ORGANISM	West Nile virus
REFERENCE	Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae; Flavivirus; Japanese encephalitis virus group. 1 (sites)
AUTHORS	Lanciotti,R.S., Roehrig,J.T., Deubel,V., Smith,J., Parker,W., Steele,K., Crise,B., Volpe,K.E., Crabtree,M.B., Scherret,J.H., Hall,R.A., MacKenzie,J.S., Cropp,C.B., Panigrahy,B., Ostlund,E., Schmitt,B.M., Malkinson,M., Banet,C., Weissman,J., Komar,N., Savage,H.M., Stone,W., McNamara,T. and Gubler,D.J.
TITLE	Origin Of the West Nile virus responsible for an outbreak of encephalitis in the northeastern United States
JOURNAL	Science 286 (5448), 2333-2337 (1999)
MEDLINE	20070288
PUBMED	10600742
REFERENCE	2 (bases 1 to 2440)
AUTHORS	Farker,M.D., Crise,B.J., Clayton,J.M. and Smith,J.F.
TITLE	Direct Submission
JOURNAL	Submitted (13-OCT-1999) Virology Division, U.S. Army Medical Research Institute of Infectious Diseases, Bldg. 1425 Fort Detrick, Frederick, Maryland 21702, USA
FEATURES source	Location/Qualifiers 1..2440 /organism="West Nile virus" /viroion
JOURNAL REFERENCE AUTHORS	from 1997 to 2000 Unpublished 2 (bases 1 to 1648) Banet,C., Brill,A., Samina,I., Yadin,H., Straum,Y., Weisman,J., Pokamonski,S., King,R., Deubel,V. and Malkinson,M.
TITLE JOURNAL	Direct Submission Submitted (01-MAY-2001) Kimron Veterinary Institute, Beit Degan 50250, Israel
FEATURES source	Location/Qualifiers 1..1648 /organism="West Nile virus" /viroion /mol_type="genomic RNA" /specific_host="white eyed gull" /db_xref="taxon:11082" <1..->1648 /codon_start=3 /product="polyprotein" /protein_id="AAL37596.1" /db_xref="GI:17226061" /translation="SLIGLKRAMLSLIDGKPIRFLVALLAFFRTAIAPTRAVLDWR RGWQTKAMHLLSPFKELGTLSAINRRSKOKRGKGKTGIAVMIGLIASVGAVTIS NFQCKVTNATVDVTIPTAGKNLCIVRAMDGYMCDDTITVECPVLSDNDP EIDWCNKTSAYYGRYCGRTKRHSRRSRSLTVQHGESLANKKGANWDSTKATRY LVKTSWLNRNGYALVAAVICGMGLSNMTGVFVLLLPAIPAYSFCNLGMSNRDF LEGVSATWDLVLDEGSCVTIMSKOKEPTIDVKMMMEANLAERYSICYLATVSDLS TKAACPTGEAHNRADPAFVRCQGVDRGNMGCGILFGKSIDTCAKFACTKAIG RTILKENIKYEVAIFVHGPTTVESHGNYSTOVGATOAGRFSITPAAPSYYTLKLGEYE VVDCPEPSGIDTIYVMTVGTFTFLVHREWFMDLNLPSSAGSTVVRNRETLMEE EPHATKQSIALGSOGEGHALAGAIPIVFESSNTVKLTSG"
CDS	Query Match 100.0%; Score 22; DB 14; Length 1648; Best Local Similarity 100.0%; Pred. No. 1.5; Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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DEFINITION	West Nile virus structural protein precursor, gene, partial cds.
ACCESSION	AF194117
VERSION	AF194117.1 GI:6715269
SOURCE	West Nile virus
ORGANISM	West Nile virus
REFERENCE	Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae; Flavivirus; Japanese encephalitis virus group. 1 (sites)
AUTHORS	Lanciotti,R.S., Roehrig,J.T., Deubel,V., Smith,J., Parker,W., Steele,K., Crise,B., Volpe,K.E., Crabtree,M.B., Scherret,J.H., Hall,R.A., MacKenzie,J.S., Cropp,C.B., Panigrahy,B., Ostlund,E., Schmitt,B.M., Malkinson,M., Banet,C., Weissman,J., Komar,N., Savage,H.M., Stone,W., McNamara,T. and Gubler,D.J.
TITLE	Origin Of the West Nile virus responsible for an outbreak of encephalitis in the northeastern United States
JOURNAL	Science 286 (5448), 2333-2337 (1999)
MEDLINE	20070288
PUBMED	10600742
REFERENCE	2 (bases 1 to 2440)
AUTHORS	Farker,M.D., Crise,B.J., Clayton,J.M. and Smith,J.F.
TITLE	Direct Submission
JOURNAL	Submitted (13-OCT-1999) Virology Division, U.S. Army Medical Research Institute of Infectious Diseases, Bldg. 1425 Fort Detrick, Frederick, Maryland 21702, USA
FEATURES source	Location/Qualifiers 1..2440 /organism="West Nile virus" /viroion

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 VERSION AY268133.1 GI:33242576
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 West Nile virus (WNV)
 Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 Flavivirus; Japanese encephalitis virus group.
 REFERENCE
 1 (bases 1 to 10989)
 Charrel,R.N., Brault,A.C., Gallian,P., Lemasson,J.-J., Murgue,B.,
 Murri,S., Pastorino,B., Zeller,H., de chesese,R., de Micco,P. and de
 Lamballerie,X.
 Evolutionary relationship between Old World West Nile virus
 strains. Evidence for viral gene flow between africa, the middle
 east, and europe
 Virology 315 (2), 381-388 (2003)
 JOURNAL
 MEDLINE 22949215
 PUBMED 14585341
 REFERENCE
 2 (bases 1 to 10989)
 de Lamballerie,X., Brault,A.C., Gallian,P., Lemasson,J., Murgue,B.,
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 Direct Submission
 Submitted (03-APR-2003) Virology, Medical University, 27 bd Jean
 Moulin, Marseille 13005, France
 JOURNAL
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ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.5;
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ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.5;
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DEFINITION
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VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
AUTHORS
TITLE
JOURNAL
FEATURES
source

AY278441
West Nile virus isolate Ast99-901, complete genome.
AY278441
AY278441.1 GI:30349729
West Nile virus (WNV)
West Nile virus
Viruses; sRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
1 (bases 1 to 10988)
Voronina,A.G., Philipov,A.G., Kinney,R.M., Samokhvalov,E.I.,
Savage,H.M., Alkhovsky,S.V., Tsychia,R., Sadykova,G.K.,
Shatalov,A.G., Usachev,E.V., Mokhonov,V.V., Butenko,A.M.,
Larichev,V.F., Gubler,D.J. and Lvov,D.K.
Analysis of a new variants of West Nile virus
Unpublished
2 (bases 1 to 10988)
Voronina,A.G., Philipov,A.G., Kinney,R.M., Samokhvalov,E.I.,
Savage,H.M., Alkhovsky,S.V., Tsychia,R., Sadykova,G.K.,
Shatalov,A.G., Usachev,E.V., Mokhonov,V.V., Butenko,A.M.,
Larichev,V.F., Gubler,D.J. and Lvov,D.K.
Direct Submission
Submitted (17-Apr-2003) Molecular Genetic, Ivanovsky Virology
Institute, Gamalei 16, Moscow 123098, Russia
Location/Qualifiers
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VERSION      AX576542.1 GI:27646162
KEYWORDS
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ORGANISM      Flavivirus sp.
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus.

REFERENCE
AUTHORS      Despres, P., Deubel, V., Guenet, J. L., Drouet, M. T., Malkinson, M. K.,
Banet, C. K., Frenkiel, M. P., Courageot, M. P., Coulibaly, F.,
Catteau, A., Flanand, M., Weber, P. and Ceccaldi, P. E.
TITLE      Neurovirulent strain of the west nile virus and applications thereof
JOURNAL
Patent: WO 02081511-A 1 17-OCT-2002;
INSTITUT PASTEUR (FR) ; Kimron Veterinary Institute (IL)
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ORIGIN
Query Match      100.0%; Score 22; DB 6; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
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Db      195 AGCCCTCTTCAGTCCAATCAAG 174

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LOCUS      AX577796      11029 bp      DNA      linear      PAT 08-JAN-2003
DEFINITION      Sequence 1 from Patent WO02081741.
ACCESSION      AX577796
VERSION      AX577796.1 GI:27647035
KEYWORDS
SOURCE
ORGANISM      Flavivirus sp.
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus.

REFERENCE
AUTHORS      Guenet, J. L., Mashimo, T., Simon-Chazottes, D., Montagutelli, X.,
Frenkiel, M. P., Despres, P., Deubel, V., Bonhomme, F. and Lucas, M.
TITLE      Use of products of genes of the 2'-5' oligoadenylate synthetase
family (oas) for screening antiviral agents and for detecting
responsiveness to flaviviridae infection
JOURNAL      Patent: WO 02081741-A 1 17-OCT-2002;
INSTITUT PASTEUR (FR) ; CENTRE NATIONAL DE LA RECHERCHE
SCIENTIFIQUE (CNRS) (FR)
FEATURES
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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGCCCTCTTCAGTCCCAATCAAG 22
Db 195 AGCCCTCTTCAGTCCCAATCAAG 174

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West Nile virus (WNV)
Virus; sRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
Shirato, K., Miyoshi, H., Goto, A., Ako, Y., Ueki, T., Kariwa, H. and
Takashima, I.
Correlation between viral envelope glycosylation and
neuroinvasiveness of the New York strain of the West Nile virus
Unpublished
2 (bases 1 to 11029)
Shirato, K., Kariwa, H. and Takashima, I.
Direct Submission
Submitted (28-JUL-2004) Kazuya Shirato, Graduate School of
Veterinary Medicine, Hokkaido University, Laboratory of Public
Health, Department of Environmental Veterinary Medicine; Kita-19
Nishi-9, Kita-ku, Sapporo, Hokkaido 060-0818, Japan
(E-mail:shirato@vetmed.hokudai.ac.jp, Tel:81-11-706-5213 (ex.5213),
Fax:81-11-706-5213)
On Jul 30, 2004 this sequence version replaced gi:50838780.
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Query Match 100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 195 AGCCCTCTTCAGTCCCAATCAAG 174
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LOCUS
DEFINITION
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cde, isolate: B-Sp.
AB185916
ACCESSION

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VERSION      AB185916.1  GI:50838782
KEYWORDS     West Nile virus (WNV)
SOURCE       West Nile virus
ORGANISM     Flavivirus; Japanese encephalitis virus group.

REFERENCE    1
AUTHORS      Shirato,K., Miyoshi,H., Goto,A., Ako,Y., Ueki,T., Kariwa,H. and
              Takashima,I.
TITLE        Correlation between viral envelope glycosylation and
              neuroinvasiveness of the New York strain of the West Nile virus
              Unpublished
JOURNAL      2 (bases 1 to 11029)
AUTHORS      Shirato,K., Kariwa,H. and Takashima,I.
TITLE        Direct Submission
JOURNAL      Submitted (28-JUL-2004) Kazuya Shirato, Graduate School of
              Veterinary Medicine, Hokkaido University, Laboratory of Public
              Health, Department of Environmental Veterinary Medicine; Kita-19
              Nishi-9, Kita-ku, Sapporo, Hokkaido 060-0818, Japan
              (E-mail:shirato@vetmed.hokudai.ac.jp, Tel:81-11-706-5213),
              Fax:81-11-706-5213)
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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCAATCAAG 22
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Db 195 AGCCCTCTTCAGTCAATCAAG 174
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RESULT 14
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LOCUS
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cds, isolate: B-LP.
ACCESSION AB185917
VERSION AB185917.1 GI:50838784
KEYWORDS West Nile virus (WNV)
SOURCE Viruses; sRNA positive-strand viruses, no DNA stage; Flaviviridae;
ORGANISM Flavivirus; Japanese encephalitis virus group.

REFERENCE 1
AUTHORS Shirato,K., Miyoshi,H., Goto,A., Ako,Y., Ueki,T., Kariwa,H. and
Takashima,I.
TITLE Correlation between viral envelope glycosylation and
neuroinvasiveness of the New York strain of the West Nile virus
Unpublished
JOURNAL 2 (bases 1 to 11029)
AUTHORS Shirato,K., Kariwa,H. and Takashima,I.
TITLE Direct Submission
JOURNAL Submitted (28-JUL-2004) Kazuya Shirato, Graduate School of
Veterinary Medicine, Hokkaido University, Laboratory of Public
Health, Department of Environmental Veterinary Medicine; Kita-19
Nishi-9, Kita-ku, Sapporo, Hokkaido 060-0818, Japan
(E-mail:shirato@vetmed.hokudai.ac.jp, Tel:81-11-706-5213),
Fax:81-11-706-5213)

FEATURES Location/Qualifiers
1..11029
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Query Match      100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. NO. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 AGCCCTCTTCAGTCCCAATCAAG 22
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Db      195 AGCCCTCTTCAGTCCCAATCAAG 174

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LOCUS      AF196835      11029 bp      linear      VRL 07-DEC-2000
DEFINITION West Nile virus strain NY99-flamingo382-99, complete genome.
ACCESSION  AF196835

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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCAAG 22
Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 16
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LOCUS West Nile virus strain NY99-eghs, complete genome.
DEFINITION
ACCESSION AF260967
VERSION AF260967.1 GI:9930133
KEYWORDS
SOURCE West Nile virus
ORGANISM West Nile virus

REFERENCE
AUTHORS
TITLE
JOURNAL

REFERENCE
AUTHORS
TITLE
JOURNAL

FEATURES
source

CDS

Location/Qualifiers

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Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
1 (bases 1 to 11029)
Bowen, M., Meyer, R.F., McKinney, N., Morrill, W. and Lanciotti, R.
Complete genomic sequence of West Nile virus equine isolate New
York 1999
Unpublished
2 (bases 1 to 11029)
Bowen, M., Meyer, R.F., McKinney, N., Morrill, W. and Lanciotti, R.
Direct Submission
Submitted (27-Apr-2000) Arbovirus Diseases Branch, Centers for
Disease Control & Prevention, Rampart Road, Fort Collins, CO 80521,
USA

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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
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Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 19

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LOCUS
DEFINITION
Accession
AF404754
VERSION
AF404754.1 GI:21929234
KEYWORDS
SOURCE
ORGANISM

West Nile virus
Viruses; sRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
1 (bases 1 to 11029)

REFERENCE
AUTHORS
Lancioti, R.S., Ebel, G.D., Deubel, V., Kerst, A.J., Murri, S.,
Meyer, R., Bowen, M., McKinney, N., Morrill, W.E., Crabtree, M.B.,
Kramer, L.D. and Roehrig, J.T.
Complete genome sequences and phylogenetic analysis of West Nile
virus strains isolated from the United States, Europe, and the
Middle East

JOURNAL
MEDLINE
PUBMED
12089180
12093177

REFERENCE
AUTHORS
Lancioti, R.S., Ebel, G.D. and Kerst, A.J.
Direct Submission
Submitted (02-AUG-2001) Division of Vector-Borne Infectious
Diseases, Centers for Disease Control & Prevention, Rampart Road,
Fort Collins, CO 80521, USA
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22089180
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REFERENCE
2 (bases 1 to 11029)
Ebel,G.D., Kerst,A.J. and Lanciotti,R.S.
Direct Submission
Submitted (02-AUG-2001) Division of Vector-Borne Infectious
Diseases, Centers for Disease Control & Prevention, Rampart Road,
Fort Collins, CO 80521, USA
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AF404756
VERSION
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KEYWORDS
West Nile virus
ORGANISM
West Nile virus

REFERENCE
1 (bases 1 to 11029)
Lanciotti, R.S., Ebel, G.D., Deubel, V., Kerst, A.J., Murri, S.,
Meyer, R., Bowen, M., McKinney, N., Morrill, W.E., Crabtree, M.B.,
Kramer, L.D. and Roehrig, J.T.
Complete genome sequences and phylogenetic analysis of West Nile
virus strains isolated from the United States, Europe, and the
Middle East
Virology 298 (1), 96-105 (2002)

JOURNAL
MEDLINE
PUBMED
12093177

REFERENCE
2 (bases 1 to 11029)
Ebel, G.D., Kerst, A.J. and Lanciotti, R.S.
Direct Submission
Submitted (02-AUG-2001) Division of Vector-Borne Infectious
Diseases, Centers for Disease Control & Prevention, Rampart Road,
Fort Collins, CO 80521, USA

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Introduction of West Nile virus in the Middle East by migrating white storks Emerging Infect. Dis. 8 (4), 392-397 (2002) 21968420 11971773 2 (bases 1 to 11029) Deubel, V., Malkinson, M. and Banet, C. Direct Submission Submitted (08-FEB-2002) CERVI, Institut Pasteur, 21 Avenue Tony Garnier, Lyon 69365, France Location/Qualifiers 1. 11029 /organism="West Nile virus" /mol_type="genomic RNA" /strain="IS-98 STD" /specific_host="sick stork" /db_xref="taxon:11082" /note="Isolated in Israel in 1998" 97..10398 /codon_start=1 /product="polyprotein precursor" /protein_id="AAL87234.1" /db_xref="GI:19387528" /translation="MSKKPGPGKSRVNNMLKRGMPRVLSLIGKRAMLSLIDGKGP RFLVALAFFRETAIAPRAVLDWRGVNKKQTAMKHLISFFKELGTLTSAINRRSGKI KRGGKTGIAVMIGLIASVGTLSNFOGKVMYVNTATDVITITPAAAGNCLCIVR ANDVGMCDDTITTECPVLSAGNDEPDCWTSKSAVYVGRCTKTHRRSRSRLT VQTHGESLANKGAMWSTKATRYLVETSWILNPGYALVAAYVGRCTKTHRRSRSRLT VFVLLMLPAYSFNCILGMSRNDPLEGSGATVVDLVLGSDCVTINSKQVPTIDVK MMNEAANLAEVRSYCLATVSDLSLTKAACPMTGEAHRNDRADPAFCRQGVVDRWG NGCGLFGKSGIDTCAKFACTKAGRTILKENIKYEVALFVHGPTTVESHGNSYQVG ATQAGRTTTPAASPITLKGEGYVTDVCPRSIGDITNAYTVYVTKTFLVHRWF MDLMLPSAGSTVWRNRETLMEFEHPHATKQSVIALGSEQALGALAGAI1PVEFSS NTVKLTSGHLKCRVKMEKQLQKGTYYGVCSKAFKFLGTDPADTGHGTVLVELOYTG PCKVPTSSVASLNDLTPVGRVLTVPNFVSVATANAKVLELEPPRGDGSVIVVGRGEQ INHHWKSSSS1GKATTTILKGAORLAALGDTANDFGSVGGFTSVGRVHVOFGGAF RLPFGMSWITKGLGALLMGINARDKSIALTFLAVGGVLLFLSVNVHADTGCAID ISQRELRCGSGVFIHNDVAMDRYKYPETPOGLAKI1QKAHKEGVCGLRSVRLSH QMEAVKDLNLTLEKNGVDLSVVEKEGMYKSAPKRLTATTEKLEIGWKAWGKSIL FAPELANNTFVVDGPETKECPTQNRANNSLEVEDFGGLTSRMLKVRKESNTTBCDS KIIGTAKNNLAIHSDLSYIERSLNDTWKLERAVLGEVKSCTWPTETHLWGDGILES DLIIPVTLAGPSNNRRPGYKTONQGPWDEGRVLDPDYCGTPTVTLSESCHGRGPA TRTTESGKLI1DNMCRSCTLPRLYTQDSCGCGWEIRFQRHDEKTLVQSOVNAYNA DMIDFQGLLVFLAATQVLRKRTAKISMPAILALVLFVGGITTVLDRYILV GAAFAESGGDVHLMALMTFKIOPVFVMSFLKARWTNOENILMLAAVFFQWYAH DARQILLWSDVLSAVAMILRAITFTTTSNVVPLALLTLCRLCLNDVIRIL LIMVGTGSI1REKRSAAKKKGASLLCLALASTGLFNPMLAAGLICDPNKRKGWPA TEVMTAVGLMFAIVGGLAELDIDSMAI1PMTIAGLMFAAFV1SGKSTDMWERTADISW ESDAEITGSSERVVRLDDGNFQMLNDPGAPWKIMLWRMCLAISATPFWALLBSV GFWITLQYTKRGGVLDTPSPKEYKKGDTTGVYRIMTRGLIGLSQAGAGVNVGCVFH TLWHTTKGAALMSGEGRDYPYGSVKEDRLCYGGPKLQHKWNGQDVQMI1VVEGPN VKNVOTKGFVKTPEGEI1GAVTLDPTTSGSPIVDKNGDVLGXNGVIMPNGSVIS AIVQGERMDEPI1PAGFEPEMLRKKQITVLDLHPGAKTRRILPQI1KEALNRLRTAV LAPTRVAAEALRGPIRYQTSAPREHNGNEI1VDVMDCHATLTHRLMSPHRPVNY NLFWDEAHFTDPASIAARGYI1STKVELGEAAA1FMTATPPTGSDPPFESNPISDLQ TEIPDRANNSGYEIT1TEYGTVMFPVSPVMGNEI1ALCLQRAKKVQALNRSKYETEY PCKNDNDWDFVIT1DISEGANFKASRVIDSRSKVPKTI1TEGEGRVILGEP5AVTAA SAAQRGRIGRNP5OVDEYCYGCHTNEDDSNFHWTKEARIMLDNINMPNGLIAQFYQ PEREKVYTMGEYRLRGERKNFLELRADLPVLAAYKVAAGSVYHNRWCDFGPR TNTILEDNNNEVEVIT1KLGERRKILRPRIDARYSDHQAALKAFDPASGKRSQIGLIEV LQKPEHFMGKTWEALDTPMYVATAEKGRAHRMALEELPDALQI1ALIALISVMTMG VFFLMQRLKGI1KGLGAVLGATVFCFMAEVPGTAKIMLLLSLMLI1VILPEPEK QRSQTDNQLAVE1K1CVMTLSVSAANEMGLDKTSIDSSLFQCRIT1EYKENSMEGFL LDLRPATWSLYAVTAVTLPLKHLITSDY1NTSLTSINTVQASALFTLARGFPFVDV GVSALLAAGCGOVTLVVMTAATLLFCHYAVMPVQMAEAMRSQARTAGI1WKNA GVYSAALLAAGCGOVTLVVMTAATLLFCHYAVMPVQMAEAMRSQARTAGI1WKNA VVDGIVATDVPLEERT1PMQKVGQIMLILVSLAAVVPVPSKTVREAGILITAAAV TLWENGASSVNNAT1AIGLCHIMRGWLSCLSI1TWTLLIKNNEKPKLKGGAQRTLGE VWKRNLNMTKEEFTRYKKEAI11EVDRSAAKHARKGNVTGHHGVSRTAKURULVER RFLPEPVGKVIDLGCGRGMYMATQKRVQYRGYTKGGPGHEEQPLQVQSOYNWITM KSGVDVFRPECCDTLLC1DGE5SSAEVEHRT1RVLEWVEDLHWRGPRFCVKVL CPYMPKVI1EKMELLQRRYGGGLVRNPLSRNSTHEMYVWSRASGVNVHS1WNMTSQVLG	AUTHORS TITLE JOURNAL MEDLINE PUBMED REFERENCE AUTHORS TITLE JOURNAL FEATURES source CDS	us-10-729-421-35.rge	Page 16
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ORIGIN

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LOCUS

DEFINITION

AF533540

VERSION

AF533540.1

KEYWORDS

SOURCE

ORGANISM

West Nile virus

West Nile virus

Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;

Flavivirus; Japanese encephalitis virus group.

1 (bases 1 to 11029)

Huang, C., Slater, B., Rudd, R., Parchuri, N., Hull, R., Dupuis, M. and

Hindenburg, A.

First Isolation of West Nile virus from a Patient with Encephalitis

in the United States

Emerging Infect. Dis. 8 (12), 1367-1371 (2002)

2 (bases 1 to 11029)

Huang, C., Slater, B., Rudd, R., Parchuri, N., Hull, R., Dupuis, M. and

Hindenburg, A.

Direct Submission

Submitted (30-JUL-2002) Wadsworth Center, New York State Department

of Health, Box 509 Albany, NY 12201-0509, USA

Location/Qualifiers

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2001"

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ORIGIN

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Best Local Similarity 100.0%; Score 22; DB 14; Length 11029;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;


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LOCUS      11057 bp      RNA      linear      VRL 15-AUG-2004
DEFINITION West Nile virus strain Sarafend, complete genome.
ACCESSION  AY688948
VERSION     AY688948.1 GI:51095221
KEYWORDS
SOURCE      West Nile virus (MNV)
ORGANISM    West Nile virus
            Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
            Flavivirus; Japanese encephalitis virus group.
REFERENCE   1 (bases 1 to 11057)
AUTHORS     Li, J., Bhuvanathan, R. and Ng, M.-L.
TITLE       Construction and characterization of an infectious West Nile
            (Sarafend) clone
JOURNAL     Unpublished
AUTHORS     2 (bases 1 to 11057)
            Li, J., Bhuvanathan, R. and Ng, M.-L.
TITLE       Direct Submission
JOURNAL     Submitted (18-JUL-2004) Microbiology, National University of
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Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGCCCTCTTCAGTCCATCAAG 22
DB 195 AGCCCTCTTCAGTCCATCAAG 174
RESULT 25
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LOCUS      1648 bp      mRNA      linear      VRL 14-MAR-2002
DEFINITION West Nile virus isolate WN_0233 polyprotein mRNA, partial cds.
ACCESSION  AF375043
VERSION     AF375043.1 GI:19421849
KEYWORDS
SOURCE      West Nile virus
ORGANISM    West Nile virus
            Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
            Flavivirus; Japanese encephalitis virus group.
REFERENCE   1 (bases 1 to 1648)
AUTHORS     Hindiye, M., Shulman, L.M., Mendelson, E., Weiss, L., Grossman, Z. and
            Bin, H.
TITLE       Isolation and characterization of West Nile virus from the blood of
            viremic patients during the 2000 outbreak in Israel
```

JOURNAL Emerging Infect. Dis. 7 (4), 748-750 (2001)
MEDLINE 21469825
PUBMED 11585544
REFERENCE 2 (bases 1 to 1648)
AUTHORS Hindiyyeh,M., Shulman,L.M., Mendelson,E., Grossman,Z., Weiss,L. and Bin,H.
TITLE Direct Submission
JOURNAL Submitted (30-APR-2001) Central Virology Laboratory, Ministry of Health, Public Health Laboratories, Sheba Medical Center, Tel Hashomer 52621, Israel
FEATURES
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ORIGIN

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Best Local Similarity 100.0%; Pred. No. 5.1;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2 GCCCTCTTCAGTCCATCAAG 22
Db 25 GCCCTCTTCAGTCCATCAAG 5

Search completed: September 6, 2005, 20:29:45
Job time : 775.688 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 16:01:23 ; Search time 735.656 Seconds
(without alignments)
1383.200 Million cell updates/sec

Title: US-10-729-421-34

Perfect score: 21

Sequence: 1 ccgggtgtcaatgctaaa 21

Scoring table:

IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 100 summaries

Database : GenEmbl.*

1: gb_ba.*

2: gb_htg.*

3: gb_in.*

4: gb_om.*

5: gb_ov.*

6: gb_pat.*

7: gb_ph.*

8: gb_pl.*

9: gb_pr.*

10: gb_ro.*

11: gb_rts.*

12: gb_sy.*

13: gb_un.*

14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	21	100.0	240	14 WNF42SAA	M32560 West Nile v
2	21	100.0	2440	14 AF194117	AF194117 West Nile
3	21	100.0	10664	14 KUNCG	D00246 Kunjin viru
4	21	100.0	10842	14 AY278442	AY278442 West Nile
5	21	100.0	10845	14 AY277252	AY277252 West Nile
6	21	100.0	10945	14 AF202541	AF202541 West Nile
7	21	100.0	10962	14 WNFCCG	M12294 West Nile v
8	21	100.0	10972	14 AF317203	AF317203 West Nile
9	21	100.0	10975	14 AF206518	AF206518 West Nile
10	21	100.0	10984	14 AY262283	AY262283 West Nile
11	21	100.0	10989	14 AY268132	AY268132 West Nile
12	21	100.0	10989	14 AY268133	AY268133 West Nile
13	21	100.0	11022	14 AY274504	AY274504 Kunjin vi
14	21	100.0	11022	14 AY274505	AY274505 Kunjin vi
15	21	100.0	11028	14 AY490240	AY490240 West Nile
16	21	100.0	11029	6 AX576542	AX576542 Sequence
17	21	100.0	11029	6 AX577796	AX577796 Sequence
18	21	100.0	11029	14 AB185914	AB185914 West Nile
19	21	100.0	11029	14 AB185915	AB185915 West Nile

20	21	100.0	11029	14 AB185916	AB185916 West Nile
21	21	100.0	11029	14 AB185917	AB185917 West Nile
22	21	100.0	11029	14 AF196835	AF196835 West Nile
23	21	100.0	11029	14 AF260967	AF260967 West Nile
24	21	100.0	11029	14 AF260968	AF260968 West Nile
25	21	100.0	11029	14 AF260969	AF260969 West Nile
26	21	100.0	11029	14 AF404753	AF404753 West Nile
27	21	100.0	11029	14 AF404754	AF404754 West Nile
28	21	100.0	11029	14 AF404755	AF404755 West Nile
29	21	100.0	11029	14 AF404756	AF404756 West Nile
30	21	100.0	11029	14 AF404757	AF404757 West Nile
31	21	100.0	11029	14 AF481864	AF481864 West Nile
32	21	100.0	11029	14 AF533540	AF533540 West Nile
33	21	100.0	11029	14 AY289214	AY289214 West Nile
34	21	100.0	11057	14 AY688948	AY688948 West Nile
35	20	95.2	10998	14 AY278441	AY278441 West Nile
36	19.4	92.4	4673	14 SLOCME	M16614 St. Louis e
37	19.4	92.4	10741	14 AY277251	AY277251 West Nile
38	18	85.7	112979	5 BX088699	BX088699 Zebrafish
39	17.8	84.8	2379	14 S75726	S75726 Japanese en
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78	17.8	84.8	10976	14 AF221499	AF221499 Japanese
79	17.8	84.8	10976	14 AF221500	AF221500 Japanese
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89	17.8	84.8	10977	14 AF080251	AF080251 Japanese
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91	17.8	84.8	11014	14 AF161266	AF161266 Murray Va
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ALIGNMENTS

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DEFINITION West Nile virus (WN) 5' terminal region of genome.
ACCESSION M32560
VERSION M32560.1 GI:336165
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REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
COMMENT
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ORIGIN
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Best Local Similarity 100.0%; Pred. No. 1.1;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGCAATATGCTAAA 21
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Db 129 CCGGGCTGCAATATGCTAAA 149

RESULT 2
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LOCUS AF194117 2440 bp RNA linear VRL 19-JAN-2000
DEFINITION West Nile virus structural protein precursor, gene, partial cds.
ACCESSION AF194117
VERSION AF194117.1 GI:6715269
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ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
COMMENT
FEATURES
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QY 1 CCGGGCTGCAATATGCTAAA 21
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Db 129 CCGGGCTGCAATATGCTAAA 149

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REFERENCE
AUTHORS Lanciotti,R.S., Roehrig,J.T., Deubel,V., Smith,J., Parker,M.,
Steele,K., Crise,B., Volpe,K.E., Crabtree,M.B., Scherret,J.H.,
Hall,R.A., MacKenzie,J.S., Cropp,C.B., Panigrahy,B., Ostlund,E.,
Schmitt,B., Malkinson,M., Banet,C., Weissman,J., Komar,N.,
Savage,H.M., Stone,W., McNamara,T. and Gubler,D.J.
TITLE Origin of the West Nile virus responsible for an outbreak of
encephalitis in the northeastern United States
JOURNAL Science 286 (5448), 2333-2337 (1999)
MEDLINE 20070288
PUBMED 10600742
REFERENCE
2 (bases 1 to 2440)
Parker,M.D., Crise,B.J., Clayton,J.M. and Smith,J.F.
Direct Submission
JOURNAL Submitted (13-OCT-1999) Virology Division, U.S. Army Medical
Research Institute of Infectious Diseases, Bldg. 1425 Fort Detrick,
Frederick, Maryland 21702, USA
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RESULT 3
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LOCUS KUNCG 10664 bp RNA linear VRL 17-FEB-1998
DEFINITION Kunjin virus gene for polyprotein (C, prM, E, NS1, NS2A, NS2B, NS3,
NS4A, NS4B, NS5), complete cds.
ACCESSION D00246
VERSION D00246.1 GI:221966
KEYWORDS M (membrane protein); prM (precursor of M); NS5; NS4B; NS4A; NS3;
NS2B; NS2A; NS1; E (envelope protein); C (core protein);
polyprotein.
SOURCE
ORGANISM Kunjin virus
Viruses; sRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
1 (bases 1 to 10664)
Coia,G., Parker,M.D., Speight,G., Byrne,M.E. and Westaway,E.G.
REFERENCE
AUTHORS
TITLE
Nucleotide and complete amino acid sequences of Kunjin virus:
definitive gene order and characteristics of the virus-specified
proteins

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J. Gen. Virol. 69 (Pt 1), 1-21 (1988)
88089524
MEDLINE
PUBMED
COMMENT

A kunjin (KUN) virus cDNA sequence of 10664 nucleotides which encoded a single open reading frame for 3433 amino acids was obtained and compared with the complete amino acid sequences of yellow fever and West Nile viruses. Partial N-terminal amino acid analyses of KUN virus-specified proteins identified the polyprotein cleavage sites and the definitive gene order. Three stop codons in the correct reading frame occur within the first 25 nucleotides beyond the 3' end of the coding sequence.

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Query Match 100.0%; Score 21; DB 14; Length 10664;
Best Local Similarity 100.0%; Pred. NO. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGCTCAATATGCTAAA 21
Db 108 CCGGGCTGCTCAATATGCTAAA 128

RESULT 4
AY278442
LOCUS AY278442 10842 bp RNA linear VRL 03-MAY-2003
DEFINITION West Nile virus isolate LEIV-VI900-27924, complete genome.
ACCESSION AY278442
VERSION AY278442.1 GI:30349731
KEYWORDS
SOURCE West Nile virus (WNV)
ORGANISM West Nile virus
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
REFERENCE 1 (bases 1 to 10842)
AUTHORS Sadykova, G.K., Prilipov, A.G., Kinney, R.M., Samokhvalov, E.I.,
Savage, H.M., Alkhovsky, S.V., Tsyhia, R., Gromashevsky, V.L.,
Usachev, E.V., Mokhnov, V.V., Voronina, A.G., Butenko, A.M.,
Larichev, V.F., Gubler, D.J. and Lvov, D.K.
Analysis of a new variants of West Nile virus
TITLE Unpublished
JOURNAL 2 (bases 1 to 10842)
REFERENCE Sadykova, G.K., Prilipov, A.G., Kinney, R.M., Samokhvalov, E.I.,
AUTHORS Savage, H.M., Alkhovsky, S.V., Tsyhia, R., Gromashevsky, V.L.,
Usachev, E.V., Mokhnov, V.V., Voronina, A.G., Butenko, A.M.,
Larichev, V.F., Gubler, D.J. and Lvov, D.K.
Direct Submission
TITLE Submitted (17-APR-2003) Molecular Genetic, Ivanovsky Virology
JOURNAL Institute, Gamalei 16, Moscow 123098, Russia
FEATURES Location/Qualifiers
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QY 1 CCGGGCTGCAATATGCTAAA 21
DB 129 CCGGGCTGCAATATGCTAAA 149

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West Nile virus isolate LEIV-Vlg99-27889, complete genome.
DEFINITION
ACCESSION AY277252
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ORIGIN

Query Match 100.0%; Score 21; DB 14; Length 10845;
 Best Local Similarity 100.0%; Pred. No. 1.6;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGCAATATGCTAAA 21
 Db 129 CCGGGCTGCAATATGCTAAA 149

RESULT 6.

AF202541
 LOCUS
 DEFINITION West Nile virus strain HNY1999 polyprotein (C, prM, E, NS1, NS2a, NS2b, NS3, NS4a, NS4b, NS5), complete cds.
 ACCESSION AF202541
 VERSION AF202541.1 GI:6581069
 KEYWORDS
 SOURCE West Nile virus
 ORGANISM
 Viruses; serNA positive-strand viruses, no DNA stage; Flaviviridae; Flavivirus; Japanese encephalitis virus group.
 REFERENCE 1 (bases 1 to 10945)
 AUTHORS Jia,X.Y., Briese,T., Jordan,I., Rambaut,A., Chi,H.C., Mackenzie,J.S., Hall,R.A., Scherret,J. and Lipkin,W.I.
 TITLE Genetic analysis of West Nile New York 1999 encephalitis virus
 JOURNAL Lancet 354 (9194), 1971-1972 (1999)
 MEDLINE 20086017
 PUBMED 10622305
 REFERENCE 2 (bases 1 to 10945)
 AUTHORS Jia,X.Y., Briese,T., Jordan,I. and Lipkin,W.I.
 TITLE Direct Submission
 JOURNAL Submitted (06-NOV-1999) Emerging Diseases Laboratory, Dept. Microbiology & Molecular Genetics and Neurology, University of California, Irvine, 3101 Gillespie Neuroscience Facility, Irvine, CA 92697-4292, USA
 FEATURES
 source Location/Qualifiers
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ORIGIN

Query Match 100.0%; Score 21; DB 14; Length 10962;
 Best Local Similarity 100.0%; Pred. No. 1.6;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGCAATATGCTAAA 21

Db 129 CCGGGCTGCAATATGCTAAA 149
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RESULT 8

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 LOCUS AF317203 10972 bp RNA linear VRL 11-FEB-2001
 DEFINITION West Nile virus VLG-4 polyprotein precursor, gene, complete cds.
 ACCESSION AF317203
 VERSION AF317203.1 GI:12744408
 KEYWORDS
 SOURCE West Nile virus
 ORGANISM West Nile virus
 Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae; Flavivirus; Japanese encephalitis virus group.
 REFERENCE 1 (bases 1 to 10972)

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

FEATURES

source

Platonov,A.E., Karan,L., Yazishina,S., Obukhov,I.L., Shipulina,O.
 and Shipulin,G.A.
 Genetic similarity of West Nile viruses caused epidemics in
 Volgograd 1999 and Romania 1996
 Unpublished
 2 (bases 1 to 10972)
 Karan,L., Yazishina,S., Obukhov,I.L., Shipulina,O., Shipulin,G.A.
 and Platonov,A.E.
 Direct Submission
 Submitted (26-OCT-2000) Central Research Institute of Epidemiology,
 Novogireevskaya Str. 3A, Moscow 111123, Russia
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ORIGIN

Query Match 100.0%; Score 21; DB 14; Length 10972;
Best Local Similarity 100.0%; Pred. No. 1.6; Indels 0; Gaps 0;
Matches 21; Conservative .0; Mismatches 0;

Qy 1 CCGGGCTGCAATATGCTAAA 21
|||||
Db 97 CCGGGCTGCAATATGCTAAA 117
|||||

RESULT 9
AF206518 10975 bp DNA linear VRL 08-MAY-2000
LOCUS West Nile virus isolate 2741, complete genome.
DEFINITION AF206518
ACCESSION AF206518
VERSION AF206518.2 GI:7717200

KEYWORDS
SOURCE West Nile virus
ORGANISM West Nile virus
Viruses; SRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.

REFERENCE 1 (bases 1 to 10975)
Anderson, J.F., Vosebrinck, T.G., and Andreadis, T.G.,
Waken, E.M., French, R.A., Garmendia, A.E. and Van Kruiningen, H.J.
Isolation of West Nile virus from mosquitoes, crows, and a Cooper's
hawk in Connecticut

JOURNAL Science 286 (5448), 2331-2333 (1999)
MEDLINE 20070287
PUBMED 10600741

REFERENCE 2 (bases 1 to 10975)
Vosebrinck, C.R., Anderson, J.F. and Andreadis, T.G.
Genome Sequence of West Nile Virus from Culex pipiens isolate
Unpublished

JOURNAL
TITLE
REFERENCE 3 (bases 1 to 10975)
Anderson, J.F., Andreadis, T.G. and Vosebrinck, C.R.
Direct Submission
Submitted (18-Nov-1999) Soil and Water, Connecticut Agricultural

REFERENCE
AUTHORS
TITLE
JOURNAL
REMARK
COMMENT
FEATURES
source

Experiment Station, 123 Huntington Street, New Haven, CT 06511, USA
4 (bases 1 to 10975)
Anderson, J.F., Andreadis, T.G. and Vosebrinck, C.R.
Direct Submission
Submitted (08-MAY-2000) Soil and Water, Connecticut Agricultural
Experiment Station, 123 Huntington Street, New Haven, CT 06511, USA
Sequence update by submitter
On May 8, 2000 this sequence version replaced gi:6636507.

Location/Qualifiers
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ORIGIN

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 Db 111 CCGGGCTGTCATATGCTAA 131

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 VERSION AY262283.1 GI:30230630
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 SOURCE West Nile virus (WNV)
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Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 Flavivirus; Japanese encephalitis virus group.
 1 (bases 1 to 10984)
 Charrel,R.N., Bault,A.C., Gallian,P., Lemasson,J.-J., Murgue,B.,
 Murri,S., Pastorino,B., Zeller,H., de Cheesse,R., de Micco,P. and de
 Lamballerie,X.
 Evolutionary relationship between Old World West Nile virus
 strains. Evidence for viral gene flow between africa, the middle
 east, and europe
 Virology 315 (2), 381-388 (2003)
 22949215
 PUBMED 14585341

2 (bases 1 to 10984)
 Bault,A.C. and de Lamballerie,X.
 Direct Submission
 TITLE Submitted (25-MAR-2003) Division of Vector-Borne Infectious
 Diseases, Centers for Disease Control and Prevention, P.O. Box
 2087, Fort Collins, CO 80522, USA

FEATURES

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100.0%; Score 21; DB 14; Length 10984;

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Charel,R.N.
 Direct Submission
 Submitted (03-APR-2003) Virology, Medical University, 27 bd Jean
 Moulin, Marseille 13005, France
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 Kunjin virus
 Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 Flavivirus; Japanese encephalitis virus group.
 1 (bases 1 to 11022)
 Liu,W.J., Chen,H.B. and Khromykh,A.A.
 Molecular and Functional Analyses of Kunjin Virus Infectious cDNA
 Clones Demonstrate the Essential Roles for NS2A in Virus Assembly
 and for a Nonconservative Residue in NS3 in RNA Replication
 J. Virol. 77 (14), 7804-7813 (2003)
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 2 (bases 1 to 11022)
 Khromykh,A.A., Liu,W.J. and Chen,H.B.
 Direct Submission
 Submitted (11-APR-2003) Clinical Medical Virology Centre,
 University of Queensland/Sir Albert Sakzewski Virus Research
 Centre, Royal Children's Hospital, Herston Rd., Herston, Brisbane,
 QLD 4029, Australia
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Query Match 100.0%; Score 21; DB 14; Length 11022;
Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 129 CCGGGCTGCAATATGCTAAA 149
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CDS

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Kunjin virus
Kunjin virus
Viruses; serNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
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Khromykh, A.A., Liu, W.J. and Chen, H.B.
Direct Submission
Submitted (11-Apr-2003) Clinical Medical Virology Centre,
University of Queensland/Sir Albert Sakzewski Virus Research
Centre, Royal Children's Hospital, Herston Rd., Herston, Brisbane,
QLD 4029, Australia
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3'UTR

ORIGIN

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LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

REMARK

COMMENT

FEATURES

source

CDS

AY490240 11028 bp RNA linear VRL 08-APR-2004
West Nile virus strain Chin-01, complete genome.

AY490240.2 GI:46277828

West Nile virus (WNV)

West Nile virus

Viruses; SARNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.

1 (bases 1 to 11028)

Jiang, T., Qin, B. and Deng, Y.

Sequence determination and analysis of West Nile Virus Chin strain
Unpublished

2 (bases 1 to 11028)

Jiang, T., Qin, B. and Deng, Y.

Direct Submission

Submitted (28-NOV-2003) Virology, Institute of Microbiology and
Epidemiology, FengTai Dongda Street, Beijing 100071, China

3 (bases 1 to 11028)

Jiang, T., Qin, B. and Deng, Y.

Direct Submission

Submitted (08-APR-2004) Virology, Institute of Microbiology and
Epidemiology, FengTai Dongda Street, Beijing 100071, China

Sequence update by submitter

On Apr 8, 2004 this sequence version replaced gi:40362614.

Location/Qualifiers

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ORIGIN

Query Match 100.0%; Score 21; DB 14; Length 11028;
Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 129 CCGGGCTGCAATATGCTAAA 149

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DEFINITION Sequence 1 from Patent WO02081511.
ACCESSION AX576542
VERSION AX576542.1 GI:27646162
KEYWORDS
SOURCE Flavivirus sp.
ORGANISM Flavivirus sp.
VIRUSES; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus.

REFERENCE 1
AUTHORS Despres, P., Deubel, V., Guenet, J. L., Drouet, M. T., Malkinson, M. K., Banet, C. K., Frenkel, M. P., Courageot, M. P., Coulbaly, F., Cateau, A., Flamand, M., Weber, P. and Ceccaldi, P. E.
TITLE Neurovirulent strain of the west nile virus and applications thereof
JOURNAL Patent: WO 02081511-A 1 17-OCT-2002;
INSTITUT PASTEUR (FR) ; Kimron Veterinary Institute (IL)

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ORIGIN

Query Match 100.0%; Score 21; DB 6; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CCGGGCTGCAATATGCTAAA 21
Db 129 CCGGGCTGCAATATGCTAAA 149

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Flavivirus sp.
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus.

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Guenet,J.L., Mashimo,T., Simon-Chazottes,D., Montagutelli,X.,
Freinkel,M.P., Despres,P., Deubel,V., Bonhomme,F. and Lucas,M.
Use of products of genes of the 2'-5' oligoadenylate synthetase
family (oas) for screening antiviral agents and for detecting
responsiveness to flaviviridae infection
Patent: WO 02081741-A 1 17-OCT-2002;
INSTITUT PASTEUR (FR) ; CENTRE NATIONAL DE LA RECHERCHE
SCIENTIFIQUE (CNRS) (FR)

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Sequence 1 from Patent WO02081741.
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Flavivirus sp.
Flavivirus sp.
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus.

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Guenet,J.L., Mashimo,T., Simon-Chazottes,D., Montagutelli,X.,
Freinkel,M.P., Despres,P., Deubel,V., Bonhomme,F. and Lucas,M.
Use of products of genes of the 2'-5' oligoadenylate synthetase
family (oas) for screening antiviral agents and for detecting
responsiveness to flaviviridae infection
Patent: WO 02081741-A 1 17-OCT-2002;
INSTITUT PASTEUR (FR) ; CENTRE NATIONAL DE LA RECHERCHE
SCIENTIFIQUE (CNRS) (FR)

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ORIGIN

Query Match 100.0%; Score 21; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21
|||||
Db 129 CCGGGCTGTCATATGCTAAA 149

RESULT 19
AB185915
LOCUS
DEFINITION West Nile virus gene for polyprotein precursor protein, complete cds isolate: 6-SP.
ACCESSION AB185915
VERSION AB185915.2 GI:50872125
KEYWORDS
SOURCE West Nile virus (WNV)
ORGANISM West Nile virus

Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae; Flavivirus; Japanese encephalitis virus group.
1 Shirato, K., Miyoshi, H., Goto, A., Ako, Y., Ueki, T., Kariwa, H. and Takashima, I.
Correlation between viral envelope glycosylation and neuroinvasiveness of the New York strain of the West Nile virus
Unpublished
2 (bases 1 to 11029)
Shirato, K., Kariwa, H. and Takashima, I.
Direct Submission
Submitted (28-JUL-2004) Kazuya Shirato, Graduate School of Veterinary Medicine, Hokkaido University, Laboratory of Public Health, Department of Environmental Veterinary Medicine; Kita-19 Nishi-9, Kita-ku, Sapporo, Hokkaido 060-0818, Japan
(E-mail: shirato@vetmed.hokudai.ac.jp, Tel:81-11-706-5213 (ex.5213), Fax:81-11-706-5213)
On Jul 30, 2004 this sequence version replaced gi:50838780.

FEATURES

Source

1..11029
/organism="West Nile virus"
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ORIGIN

Query Match 100.0%; Score 21; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGCAATATGCTAAA 21
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Db 129 CCGGGCTGCAATATGCTAAA 149

RESULT 20

AB185916 11029 bp RNA linear VRL 30-JUL-2004
LOCUS
DEFINITION
West Nile virus gene for polyprotein precursor protein, complete
cde, isolate: B-SP.

ACCESSION
AB185916 1 GI:50838782

VERSION

KEYWORDS

SOURCE

ORGANISM

West Nile virus (WNV)

Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;

Flavivirus; Japanese encephalitis virus group.

REFERENCE

AUTHORS

Shirato, K., Miyoshi, H., Goto, A., Ako, Y., Ueki, T., Kariwa, H. and

Takahashina, I.

Correlation between viral envelope glycosylation and

neuroinvasiveness of the New York strain of the West Nile virus

unpublished

2 (bases 1 to 11029)

Shirato, K., Kariwa, H. and Takashima, I.

Submitted (28-JUL-2004) Kazuya Shirato, Graduate School of

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Nishi-9, Kita-ku, Sapporo, Hokkaido 060-0818, Japan

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Fax: 81-11-706-5213)

Location/Qualifiers

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ORIGIN

Query Match 100.0%; Score 21; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGCAATATGCTAAA 21
|||||
Db 129 CCGGGCTGCAATATGCTAAA 149

RESULT 21

AB185917 11029 bp RNA linear VRL 30-JUL-2004
LOCUS
DEFINITION
West Nile virus gene for polyprotein precursor protein, complete
cde, isolate: B-UP.
ACCESSION
AB185917
VERSION
AB185917.1 GI:50838784
KEYWORDS

SOURCE West Nile virus (WNV)
ORGANISM West Nile virus
REFERENCE Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
AUTHORS Flavivirus; Japanese encephalitis virus group.

1 Shirato,K., Miyoshi,H., Goto,A., Ako,Y., Ueki,T., Kariwa,H. and Takashima,I.
Correlation between viral envelope glycosylation and neuroinvasiveness of the New York strain of the West Nile virus Unpublished

2 (bases 1 to 11029)
Shirato,K., Kariwa,H. and Takashima,I.
Direct Submission
Submitted (28-JUL-2004) Kazuya Shirato, Graduate School of Veterinary Medicine, Hokkaido University, Laboratory of Public Health, Department of Environmental Veterinary Medicine; Kita-19 Nishi-9, Kita-Ku, Sapporo, Hokkaido 060-0818, Japan
(E-mail:shirato@vetmed.hokudai.ac.jp, Tel:81-11-706-5213 (ex.5213), Fax:81-11-706-5213)

FEATURES
source Location/Qualifiers
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Query Match 100.0%; Score 21; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DB 129 CCGGGCTGTCATATGCTAAA 149
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ACCESSION AF196835
VERSION AF196835.2 GI:11597239
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SOURCE West Nile virus
ORGANISM West Nile virus
REFERENCE Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
AUTHORS Flavivirus; Japanese encephalitis virus group.
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Lancioti,R.S., Roehrig,J.T., Deubel,V., Smith,J., Parker,M.,
Steele,K., Crise,B., Volpe,K.E., Crabtree,M.B., Scherret,J.H.,
Hall,R.A., MacKenzie,J.S., Cropp,C.B., Panigrahy,B., Oestlund,E.,
Schmitt,B., Malkinson,M., Banet,C., Weisman,J., Komar,N.,
Savage,H.M., Stone,W., McNamara,T. and Gubler,D.J.
Origin of the West Nile virus responsible for an outbreak of
encephalitis in the northeastern United States
Science 286 (5448), 2333-2337 (1999)
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PUBMED 10600742
REFERENCE 2 (bases 1 to 11029)
Lancioti,R., Roehrig,J., Volpe,K. and Panigrahy,B.
Direct Submission
Submitted (20-OCT-1999) Division of Vector-Borne Diseases, Centers
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80521, USA
3 (bases 1 to 11029)
Lancioti,R., Roehrig,J., Volpe,K. and Panigrahy,B.
Direct Submission
Submitted (07-DEC-2000) Division of Vector-Borne Diseases, Centers
for Disease Control and Prevention, Rampart Road, Fort Collins, CO
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Sequence update by submitter
On Dec 7, 2000 this sequence version replaced gi:6636174.
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Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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DB 129 CCGGGCTGTCATATGCTAAA 149
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LOCUS DEFINITION
AF260968 West Nile virus strain Egl01, complete genome.
VERSION AF260968.1 GI:9930135
KEYWORDS
SOURCE
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REFERENCE
AUTHORS Bowen, M., Meyer, R.F., McKinney, N., Morrill, W. and Lanciotti, R.
TITLE Complete genomic sequence of West Nile virus strain Egl01
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 11029)
AUTHORS Bowen, M., Meyer, R.F., McKinney, N., Morrill, W. and Lanciotti, R.
TITLE Direct Submission
JOURNAL Submitted (27-Apr-2000) Arbovirus Diseases Branch, Centers for Disease Control & Prevention, Rampart Road, Fort Collins, CO 80521, USA

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ORIGIN

Query Match 100.0%; Score 21; DB 14; Length 11029;
 Best Local Similarity 100.0%; Pred. No. 1.6;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGTCATATGCTAAA 21
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 Db 129 CCGGGCTGTCATATGCTAAA 149

RESULT 25
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 ORGANISM West Nile virus
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 Flavivirus; Japanese encephalitis virus group.
 REFERENCE 1 (bases 1 to 11029)
 AUTHORS Savage, H.M., Celano, C., Nicolescu, G., Karabatsos, N., Lanciotti, R.,

TITLE
 JOURNAL
 MEDLINE
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 REFERENCE
 AUTHORS
 TITLE
 JOURNAL

Vladimirescu, A., Laiv, L., Ungureanu, A., Romanca, C. and Tsai, T.F.
 Entomologic and avian investigations of an epidemic of West Nile
 fever in Romania in 1996, with serologic and molecular
 characterization of a virus isolate from mosquitoes
 Am. J. Trop. Med. Hyg. 61 (4), 600-611 (1999)
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 2 (bases 1 to 11029)
 Bowen, M., Meyer, R.F., McKinney, N., Morrill, W. and Lanciotti, R.
 Direct Submission
 Submitted (27-APR-2000) Arbovirus Diseases Branch, Centers for
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 USA

FEATURES
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 ANDVGYMCDTITYECPVLSAGNDPEDIDCWCTKSAYVYGRCTKTRHSRRSRSLT
 VOTHGESTLANKKAGMWDSTKATRYLVKTESWILRNPYALVAACVGMWLSRSMQV
 VFVLLLVAPAVSFNCLGMSNRDPLEGVSATVWDLVLEGSCTVIMSCKDPTDVK
 MNMERANLAEVRSYCLATVSDLSKAACTPMGEAHNDKPADPAFCVQGVVDGNG
 NCGFLPGKGSIDTCAKFACTKATGNTILKENIKYEVAIFVHGPTTVESHGYNSTQIG
 ATOAGRSITPAAPSYRTLLKGEVETVDCPSRSGIDTNAIYVMTVGTITFLVHRWF
 MDLNPWSSAGTSVWRNETLMEFEPEPHATKQSVIALGSEGHALQALAAVPEFS
 NTKLTSGHLKCRVMEKQLKGTTYGVSATKAFKFLGTPADTGHGTWVLEQVTDG
 PKCVPTSSVASLNDLTPVGRVTVNPFVSVATANAKVLIIELEPPDGSIVLVGRGEQ
 INHHWKGSSIGKATFTTLKGAORLAALGSDTAMDPSGVGVFTSVGKAHVQVGAF
 RSLFGMSWITQGLLALLGMLNADRISALTFLAVGGVLLFLSNVHADTGGCALD
 ISRQLRSGGVFIHNDVEAMWDRYKYPETPGAKIIOKAHKGVSCHGVSVRLRH
 QMWESVDELNTLKGVDLSSVVEKQMGYSKAPKRLTATTKELEIGHWAKGKSIL
 PAPELANNTFVVDGPETKECTONRAWNSLEVEDFGFGLTSTRMFLKVRNESNTGDS
 KIIGTAKNLAHSDLSYMIERSNDTWKBRVAVLGEVSKSTPETHTLWNGGILES
 DLIIPVTLAGPSNNRRPGYKTONQCPWDEGRVEIDFYCPGTTVTLSESCGRGA
 TTTTESGKLTIDWCCSCTLPRLTYQDSGCWMEIPQRHDEKTLVPSQVNAFNA
 DMIDPFGLLVFLATQEVLRKWTAKISMPAILLALLVFGGITYTDVLRVILV
 GAFAESNSGSDVHLALMATFKIQPFVNVASFVKARMTNQENILMLAAVFPQAYH
 DARQILLWEIPDVLNSLAVAMMLRAITFTTTSNVVPLALLTPGLRCLNLDVYRIL
 LAMVGISLIREKRSAAKKKASILLCLALASTGLPNMILAAGLIACDPNRKRGWA
 TEVMTAVGLMFAIVGLAELDDISMAIEMTIAGLMFAAFVISGKSDMTMIERTADLSW
 ESDAEITGSSERVDRLDDGPNQMDPGAPWKMLRMVCLVASAYTAPWALLPSVH
 GFWITLTQYTKRGVLDWTPSPKEYKGGDTTGYVIMTRGLLGSYQAGVAGWVGFVH
 TLWHTTKGAALMSGGRDLPYWGSKEDRLCGPKWKLQHKWNGQDEQVIMVVEPGN
 VKNVOTKGVFKTPEGEIAGVLDPTGTSGSPIVDKNGDVLGNGYIMPNGSVIS
 AIVQGRWDEPIPAGEPEMLRKKOITVLDLHPGAGKTRILLPOIKAEINRRLTAV
 LATRVAAEMAEALRGPIRYQTSAPREHNGNEIVDMCHALTFLHMSPHRVNY
 NLVFMDEAFTDPASIAJLSTKVELGEAAAIEMTATPGTSDPSPNSPSISOLQ
 TEIPDRWNSGYEITETIKTVMFVPSVMGNEIALCLOKAGKVVOLNRKSYETEY
 PKCKNDWDVFTITDISEMGANFKASVIDRSKSVKPTIITEGGRVILGEPSAVTAA
 SAQRGRRIGRNSQVDEYCYGHTNEDDSNFAMHTEARIMLDNINMNGLIQAQYQ
 PEREKVTIOMGEYRLRGERKNFLERLTADLPVLAAYKVAAGVSHDRWCDFGPR
 TNLTDONNEVEITKLGKRIKILRPWIDARVYSDHOALKAFKDFASGRKSQGLILEV
 LGMPEHFMKTEWALDTMYVATAEKGGRHMALEELPDALQTLIALIALLSVMTMG
 VFELLMQRKGIKIGLVGVVAVATFCFMAEVPYKTIAGMLLSLLMLVILPEEK
 QRSOTDNLQALAVLICVTLVSAVAANEMGMDTKNDISSLFGORIEAKENFSGBEFL
 LDRPATWLSYAVTTAVTLPLKHLITSYINTSLTSINVOASALTARNGFPFVDV
 GVSALLAAGCGOVTITVTVAATLFCFYAVMPGQAEAMRSQRTTAAGIMAKNA
 VVDIGVATDPELERTTPIKOKKQIMLILVSLAAVVVNPVSKTVREAGILITAAAV
 TLEMNGASSVYNAATTAIGLCHIRGGWLSCLSTIWTLLKMKPKGLKGGKAGRTLGE
 WKERLNQMTKESEFTRKKEAIIIEVRSAAKHARKSGNVGCHPVSRGTHAKRLVVER
 RFLPEVGVKIDLGGRGWCYIMATQKRVQVEYTKGPGHEEPOLVOSYGNWIVTM
 KSGVDVFESECCDILLDCIGSSSAEVEEHTIRVLEMEVDLHGRGKFCVKVLG
 CPMYKVIEMLEQLQRYGGLVRNPLRSNTHMYVWSASGVNHSVNMITSQVLLG
 RMEKRTWKQYEDVEDNLGSTRVAGKPLNSDTSKIKNRIERLREYSSTWHDENH
 PYRTWNYHGSYDVKPTGSSASLIANGVVRLLSKPDRTITNTWTAMTDITPFGOORVPK
 EKVDTKAPEPPGKVVYLNETHWMAFLAREKPRMCSREEFIKKNSNAALGAMFE
 EQNMRSAEAVDPEKFMWVDEBERAHLRGECHTCIYNMNGKREKKGFGKAGKSR
 AIWFMILGARFLFEALGFLEDHWRGNKSGGVEGLGKLGYILREYVTRPGGI
 YADDTAGWDRITRADDLENAKLELLDGEHRLARAIIEITYRHVKVVMRPAADGR
 TVMDVISRQDQSGOVVYALNTFTNLAQLVRMMEGCVIGPDDVEKLTCKGKPKV
 RTWLPENGERLSERLMAVSGDDCVKPLDDRPAFSLHPLNAMSVKRDIQEWKSTGY
 DWQVPPCSNHFTELIMKQRTLVVPCRGODELVGRARISPGAGMNVDRDTACLAKSYA
 QMWLLYFHRDRILRLMANAICSAVPVNMVPTGRTWSIHAGGEMMTTEDLVEYNRVY
 IBEENWEMEDTPEVKMSDVPYSGKREDIMCGSLIGTRTRATWAENIQAIVNQVRAIIG
 DEKYDVMSSLKRYEDTTLVEDTVL"

CDS

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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 16:01:23 ; Search time 189.656 Seconds
(without alignments)
655.473 Million cell updates/sec

Title: US-10-729-421-34
Perfect score: 21
Sequence: 1 ccgggtgtcaatgtctaaa 21

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 100 summaries

Database : N_Geneseq_16Dec04:*

1: Geneseqn1980s:*

2: Geneseqn1990s:*

3: Geneseqn2000s:*

4: Geneseqn2001as:*

5: Geneseqn2001bs:*

6: Geneseqn2002as:*

7: Geneseqn2002bs:*

8: Geneseqn2003as:*

9: Geneseqn2003bs:*

10: Geneseqn2003cs:*

11: Geneseqn2003ds:*

12: Geneseqn2004as:*

13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	21	100.0	21	12	ADQ30664
2	21	100.0	37	12	ADN36773
3	21	100.0	65	12	ADN36771
4	21	100.0	365	6	ABK51710
5	21	100.0	366	8	ABQ76684
6	21	100.0	967	12	ADQ30647
7	21	100.0	10945	13	ADR32078
8	21	100.0	10945	13	ADR67768
9	21	100.0	10962	12	ADK13681
10	21	100.0	10975	12	ADN98022
11	21	100.0	11029	8	ABZ68481
12	21	100.0	11029	10	ABV74821
13	21	100.0	11029	12	ADN98023
14	19	90.5	21	12	ADN36695
15	19	90.5	24	12	ADN36823
16	19	90.5	48	12	ADN36707
17	19	90.5	69	12	ADN36694
18	18	85.7	20	12	ADN36696
19	18	85.7	47	12	ADN36708
20	17.8	84.8	4512	2	AAQ22767

21	17.8	84.8	10818	12	ADO07431	Ado07431 Japanese
22	17.8	84.8	10968	12	ADO07437	Ado07437 Japanese
23	17.8	84.8	10976	3	ABL50890	Ab150890 Japanese
24	17.8	84.8	18563	12	ADO07466	Ado07466 Japanese
25	17.8	84.8	18563	12	ADO07465	Ado07465 Japanese
26	17.8	84.8	18565	12	ADO07467	Ado07467 Japanese
27	17.8	84.8	19038	12	ADO07468	Ado07468 Japanese
28	17.8	84.8	19038	12	ADO07469	Ado07469 Japanese
29	17.8	84.8	19040	12	ADO07470	Ado07470 Japanese
30	17	81.0	17	6	ACN01436	Acn01436 WNV Inozy
31	17	81.0	17	6	ACN00029	Acn00029 WNV Hamme
32	17	81.0	17	6	ACN09480	Acn09480 WNV minus
33	17	81.0	17	6	ACN15271	Acn15271 WNV minus
34	17	81.0	17	6	ACN14165	Acn14165 WNV minus
35	17	81.0	17	6	ACN04724	Acn04724 WNV DNazY
36	17	81.0	17	6	ACN09479	Acn09479 WNV minus
37	17	81.0	20	12	ADN36776	Adn36776 West Nile
38	16.8	80.0	2262	8	ACA54320	Aca54320 Prokaryot
39	16.4	78.1	356	9	ACH31834	Ach31834 Human bon
40	16.2	77.1	326	6	ABN75146	Abn75146 Human ORF
41	16.2	77.1	1149	10	ADB46059	Adb46059 rapC DNA
42	16.2	77.1	2023	10	ADB69062	Adb69062 C. neofor
43	16.2	77.1	2232	11	ACH98107	Ach98107 Klebsiell
44	16.2	77.1	3177	4	ABL16411	Ab116411 Drosophil
45	16.2	77.1	7018	4	ABL16410	Ab116410 Drosophil
46	16	76.2	17	6	ACN00030	Acn00030 WNV Hamme
47	16	76.2	17	6	ACN03477	Acn03477 WNV Zinzy
48	16	76.2	2005	4	AAE81805	Aaf81805 Human sec
49	16	76.2	149671	6	ABK84797	Abk84797 Human CDN
50	16	76.2	149671	9	ADB70361	Adb70361 Moesin CD
51	16	76.2	149671	12	ADJ371140	Adj371140 Human mal
52	15.8	75.2	624	6	ABN73108	Abn73108 Bovine em
53	15.8	75.2	1197	5	AAE81962	Aas81962 DNA encod
54	15.8	75.2	1208	5	AAE87482	Aas87482 DNA encod
55	15.8	75.2	1208	5	AAE93301	Aas93301 DNA encod
56	15.8	75.2	1208	5	AAE77346	Aas77346 DNA encod
57	15.8	75.2	2373	5	AAE86894	Aas86894 DNA encod
58	15.8	75.2	11184	12	ADP86274	Adp86274 Hepatitis
59	15.8	75.2	11184	12	ADP86276	Adp86276 Hepatitis
60	15.8	75.2	11184	12	ADP86277	Adp86277 Hepatitis
61	15.8	75.2	11313	12	ADP86273	Adp86273 Hepatitis
62	15.8	75.2	11313	12	ADP86264	Adp86264 Hepatitis
63	15.8	75.2	11313	12	ADP86266	Adp86266 Hepatitis
64	15.8	75.2	11313	12	ADP86265	Adp86265 Hepatitis
65	15.8	75.2	11313	12	ADP86268	Adp86268 Hepatitis
66	15.8	75.2	11313	12	ADP86270	Adp86270 Hepatitis
67	15.8	75.2	11313	12	ADP86271	Adp86271 Hepatitis
68	15.8	75.2	11313	12	ADP86272	Adp86272 Hepatitis
69	15.8	75.2	11313	12	ADP86269	Adp86269 Hepatitis
70	15.8	75.2	11313	12	ADP86275	Adp86275 Hepatitis
71	15.8	75.2	11313	12	ADP86267	Adp86267 Hepatitis
72	15.8	75.2	12306	10	ADI41414	Adi41414 BB7 nucle
73	15.8	75.2	12315	10	ADI41413	Adi41413 BB7M4RLU
74	15.8	75.2	12980	2	AAV59364	Aav59364 Hepatitis
75	15.8	75.2	12980	6	ABK87286	Abk87286 Hepatitis
76	15.8	75.2	12980	8	ACA62469	Aca62469 DNA encod
77	15.8	75.2	15065	3	AAZ36195	Aaz36195 Nucleotid
78	15.8	75.2	16847	12	ADO07464	Ado07464 Japanese
79	15.4	73.3	444	9	ACH24983	Ach24983 Human adu
80	15.4	73.3	1119	6	AAJ31757	Aaj31757 Soybean H
81	15.4	73.3	73583	12	ADQ59187	Adq59187 MSI-H car
82	15.4	73.3	222930	6	ABK84349	Abk84349 Human CDN
83	15.4	73.3	295096	11	ACN40068	Acn40068 Mouse gen
84	15.2	72.4	26	12	ADN36839	Adn36839 West Nile
85	15.2	72.4	445	4	AAE36425	Aae36425 Human car
86	15.2	72.4	445	10	ADE47119	Ade47119 Human car
87	15.2	72.4	445	13	ADJ08537	Adj08537 Human car
88	15.2	72.4	452	3	AAE82296	Aae82296 N. mening
89	15.2	72.4	914	5	AAE66345	Aae66345 DNA encod
90	15.2	72.4	1038	3	AAZ60397	Aaz60397 A dialcyl
91	15.2	72.4	1065	12	ADP72956	Adp72956 Renal tox
92	15.2	72.4	1121	2	AAQ21554	Aaq21554 Polyfunct
93	15.2	72.4	1121	6	ABK63744	Abk63744 Rat seque

94 15.2 72.4 1121 10 ADB58300 Toxicity-
 95 15.2 72.4 1121 10 ADB52851 Primary r
 96 15.2 72.4 1121 10 ADB41990 Toxicity
 97 15.2 72.4 1167 10 ADF01035 Bacterial
 C 98 15.2 72.4 1302 4 AAF58403 Rat rOCIL
 C 99 15.2 72.4 1389 11 ACH96542 Klebsiell
 100 15.2 72.4 1418 6 ABZ15428 Arabidops

ALIGNMENTS

RESULT 1
 ADQ30664
 ID ADQ30664 standard; DNA; 21 BP.
 XX
 AC ADQ30664;
 XX
 DT 23-SEP-2004 (first entry)
 XX
 DE West Nile Virus capsid gene sense primer WNVVAL.
 XX
 KW ss; primer; West Nile Virus; diagnosis.
 XX
 OS West Nile virus.
 XX
 PN WO2004055159-A2.
 XX
 PD 01-JUL-2004.
 XX
 PF 05-DEC-2003; 2003WO-US038750.
 XX
 PR 12-DEC-2002; 2002US-0432850P.
 PR 20-JUN-2003; 2003US-0480431P.
 XX
 PA (CHIR) CHIRON CORP.
 XX
 PI Shyamala V;
 XX
 DR WPI; 2004-488058/46.
 XX

XX New isolated oligonucleotides for accurately diagnosing West Nile virus
 PT infection or for capturing, detecting and quantitating West Nile virus in
 PT blood samples.
 XX
 PS Claim 1; SEQ ID NO 34; 56pp; English.
 XX
 CC The invention relates to an isolated oligonucleotide not more than 60
 CC nucleotides in length comprising a nucleotide sequence (S1) of at least
 CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
 CC 20, 21 or 23 bp) given in the specification derived from the West Nile
 CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
 CC identity to the nucleotide sequence of (S1), or complements of (S1) and
 CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
 CC end and/or the 3'-end. The detectable label is a fluorescent label
 CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
 CC (TAMRA), and 2',4',5',7',-tetrachloro-4-7-dichlorofluorescein (TER). The
 CC composition and methods are useful for accurately diagnosing West Nile
 CC virus infection or for capturing, detecting and quantitating West Nile
 CC virus in biological samples, particularly blood samples. This sequence
 CC corresponds to a PCR primer to amplify a fragment of the capsid gene of
 CC the WNV genome. The fragment is detected using the oligonucleotides of
 CC the invention.

XX Sequence 21 BP; 6 A; 5 C; 5 G; 5 T; 0 U; 0 Other;
 Query Match 100.0%; Score 21; DB 12; Length 21;
 Best Local Similarity 100.0%; Pred. No. 0.38; Mismatches 0; Gaps 0;
 Matches 21; Conservative 0; Indels 0; Indels 0; Gaps 0;
 OY 1 CCGGGCTGTCAATATGCTAAA 21
 DB 1 CCGGGCTGTCAATATGCTAAA 21

RESULT 2
 ADN36773
 ID ADN36773 standard; DNA; 37 BP.
 XX
 AC ADN36773;
 XX
 DT 15-JUL-2004 (first entry)
 XX
 DE West Nile virus detection-related oligonucleotide probe SeqID95.
 XX
 KW hybridisation assay probe; nucleic acid detection;
 KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 KW high throughput screening; probe; ss.
 XX
 OS West Nile virus.
 XX
 PN WO2004036190-A2.
 XX
 PD 29-APR-2004.
 XX
 PF 10-OCT-2003; 2003WO-US033639.
 XX
 PR 16-OCT-2002; 2002US-0418891P.
 PR 25-NOV-2002; 2002US-0429006P.
 PR 24-FEB-2003; 2003US-0449810P.
 XX
 PA (GENP-) GEN-PROBE INC.
 XX
 PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
 XX
 DR WPI; 2004-389590/36.
 XX

XX New hybridization assay probe comprising target-complementary sequence of
 PT bases, useful in detecting flavivirus, e.g. West Nile virus.
 XX
 PS Claim 55; SEQ ID NO 95; 135pp; English.
 XX
 CC This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of an oligonucleotide probe which is related
 CC to the invention.
 XX
 SQ Sequence 37 BP; 10 A; 10 C; 11 G; 6 T; 0 U; 0 Other;
 Query Match 100.0%; Score 21; DB 12; Length 37;
 Best Local Similarity 100.0%; Pred. No. 0.41;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 CCGGGCTGTCAATATGCTAAA 21
 DB 4 CCGGGCTGTCAATATGCTAAA 24

RESULT 3
 ADN36771
 ID ADN36771 standard; DNA; 65 BP.
 XX
 AC ADN36771;
 XX
 DT 15-JUL-2004 (first entry)
 XX

DE West Nile virus detection-related oligonucleotide probe SeqID93.
 XX hybridisation assay probe; nucleic acid detection;
 KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 KW high throughput screening; probe; ss.

XX West Nile virus.

OS WO2004036190-A2.

PN 29-APR-2004.

PD 10-OCT-2003; 2003WO-US033639.

PF 16-OCT-2002; 2002US-0418891P.

PR 25-NOV-2002; 2002US-0429006P.

PP 24-FEB-2003; 2003US-0449810P.

PX (GENP-) GEN-PROBE INC.

PA Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
 PI WPI; 2004-389590/36.

DR New hybridization assay probe comprising target-complementary sequence of
 XX bases, useful in detecting flavivirus, e.g. West Nile virus.

PS Example 6; SEQ ID NO 93; 135pp; English.

XX This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of an oligonucleotide probe which is related
 CC to the invention.

SQ Sequence 65 BP; 19 A; 17 C; 20 G; 9 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 12; Length 65;

Best Local Similarity 100.0%; Pred. No. 0.44; Length 65;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGTCAATATGCTAAA 21

DB 32 CCGGGCTGTCAATATGCTAAA 52

RESULT 4

ABK51710

ID ABK51710 standard; cDNA; 365 BP.

XX ABK51710;

AC 27-AUG-2002 (first entry)

DT Partial cDNA for west nile virus capsid protein.

DE Human; ss; IgE leader sequence; west nile virus capsid protein;
 KW RNA secondary structure; free energy; gene therapy; cancer;
 KW hyperproliferative disease; autoimmune disease; rheumatoid arthritis;
 KW multiple sclerosis; Sjogren's syndrome; sarcoidosis; scleroderma;
 KW insulin-dependent diabetes mellitus; autoimmune thyroiditis; psoriasis;
 KW reactive arthritis; ankylosing spondylitis; polymyositis; vasculitis;
 KW dermatomyositis; Crohn's disease; ulcerative colitis.

XX

OS West Nile virus.

XX WO200229088-A2.

PN 11-APR-2002.

PD 04-OCT-2001; 2001WO-US031451.

PF 04-OCT-2000; 2000US-0237885P.

PP (UYPE-) UNIV PENNSYLVANIA.

PX Weiner DB, Yang J;

PI WPI; 2002-416682/44.

DR Producing recombinant protein for preparing pharmaceutical compounds to
 XX treat, e.g., cancers or autoimmune disorders, comprises predicting
 XX secondary structure (SS) of mRNA and modifying DNA to give mRNA with SS
 XX having increased free energy.

PS Example 2; Fig 1; 48pp; English.

XX The invention relates to producing (M1) a protein (I) in a recombinant
 CC expression system (II) comprising: (a) predicting the secondary structure
 CC of mRNA; (b) modifying the native heterologous DNA sequence where the
 CC mRNA transcribed from the modified DNA has a secondary structure with
 CC increased free energy; and (c) using the modified DNA in (II) for
 CC production of (I). Also included are (1) an injectable pharmaceutical
 CC composition comprising a nucleic acid molecule that includes a modified
 CC coding sequence (IV) encoding a protein operably linked to regulatory
 CC elements, where (IV) comprises a higher AT or AU content relative to the
 CC AT or AU content of the native coding sequence and further comprising a
 CC pharmaceutical carrier and (2) a recombinant viral vector comprising a
 CC nucleic acid molecule that includes (IV). The method is used for
 CC producing a protein in a recombinant expression system. Use of a nucleic
 CC acid or recombinant viral vector that have modified DNA sequences to
 CC improve protein production can be used in gene therapy and for the
 CC treatment of cancers, hyperproliferative diseases, and autoimmune
 CC diseases such as rheumatoid arthritis, multiple sclerosis, Sjogren's
 CC syndrome, sarcoidosis, insulin-dependent diabetes mellitus, autoimmu
 CC thyroiditis, reactive arthritis, ankylosing spondylitis, scleroderma,
 CC polymyositis, dermatomyositis, psoriasis, vasculitis, Crohn's disease and
 CC ulcerative colitis. The present sequence is a cDNA for West Nile virus
 CC capsid protein. Fusion constructs of modified mRNA for the capsid protein
 CC and human IgE leader sequence are used in an experiment to minimise the
 CC free energy of the capsid protein mRNA. Note: The present sequence is not
 CC shown in the specification but was created using the information in
 CC figure 1 and the sequence appearing as ABK51708

SQ Sequence 365 BP; 103 A; 80 C; 109 G; 73 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 6; Length 365;

Best Local Similarity 100.0%; Pred. No. 0.57; Length 365;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGTCAATATGCTAAA 21

DB 29 CCGGGCTGTCAATATGCTAAA 49

RESULT 5

ABQ76684

ID ABQ76684 standard; DNA; 366 BP.

XX ABQ76684;

AC 13-MAY-2003 (first entry)

DT WNVcwt DNA fragment.

DE Capsid protein; WNVcwt; mRNA secondary structure; cancer;
 KW immunosuppressive; antirheumatic; cytostatic; antiulcer; neuroprotective;

KW antiarthritic; antidiabetic; antithyroid; antipsoriatic; antiviral; gene;
 KW antiparasitic; antiallergic; gene therapy; allergen; multiple sclerosis;
 KW protective immune response; hyperproliferative cell; ulcerative colitis;
 KW hyperproliferative disease; psoriasis; autoimmune disease; psoriasis;
 KW rheumatoid arthritis; Sjogren's syndrome; autoimmune thyroiditis;
 KW insulin dependent diabetes mellitus; Crohn's disease; ds.

OS West Nile virus.

XX Key Location/Qualifiers
 FT CDS 1..366
 FT /*tag= a
 FT /product= "WNVCwt"
 FT /note= "no start or stop codon given"

XX US2002123099-A1.

XX 05-SEP-2002.

XX 04-OCT-2001; 2001US-00971806.

XX 04-OCT-2000; 2000US-0237885P.

XX (WEIN/) WEINER D B.
 XX (YANG/) YANG J.

XX Weiner DB, Yang J;

XX WPI; 2003-066795/06.
 XX P-PSDB; ABG73556.

XX Producing protein in recombinant expression system involves predicting
 PT secondary structure of RNA encoding a protein and increasing free energy
 PT for the secondary structure by modifying sequence of DNA encoding the
 PT RNA.

XX Example 2; Fig 1; 25pp; English.

XX This invention describes a novel method for producing a protein by
 CC translation of mRNA from heterologous DNA sequences. The method involves
 CC predicting the secondary structure of mRNA transcribed from a native
 CC heterologous DNA sequence, modifying the sequence where mRNA transcribed
 CC from the modified DNA sequence has a secondary structure with increased
 CC free energy compared to mRNA transcribed from native DNA and using
 CC modified heterologous DNA for protein production. The products of the
 CC invention have immunosuppressive, antirheumatic, cytostatic, antitumor,
 CC neuroprotective, antiarthritic, antidiabetic, antithyroid, antipsoriatic,
 CC antiviral, antiparasitic and antiallergic activity and can be used for
 CC gene therapy. The method described is useful for producing a protein in a
 CC recombinant expression system, preferably a cell free in vitro
 CC transcription and translation system, an in vitro cell expression system,
 CC a DNA construct used in direct DNA injection, or a recombinant vector for
 CC delivery of DNA to an individual. The products of the invention are
 CC useful for eliciting broad immune responses against a target protein,
 CC i.e. proteins specifically associated with pathogens such as viruses,
 CC parasites, allergens, or the individual's own abnormal cells.
 CC Compositions containing the products of the invention confer a broad
 CC based protective immune response against hyperproliferative cells that
 CC are characteristic in hyperproliferative diseases including all forms of
 CC cancer and psoriasis. Such compositions are also useful for treating
 CC individuals suffering from autoimmune diseases including rheumatoid
 CC arthritis, multiple sclerosis, Sjogren's syndrome, insulin dependent
 CC diabetes mellitus, autoimmune thyroiditis, Crohn's disease, ulcerative
 CC colitis and psoriasis. This sequence encodes the West Nile virus wild-
 CC type capsid protein described as WNVCwt in the disclosure of the
 CC invention

XX Sequence 366 BP; 103 A; 81 C; 108 G; 74 T; 0 U; 0 Other;

XX Query Match 100.0%; Score 21; DB 8; Length 366;
 XX Best Local Similarity 100.0%; Pred. No. 0.57;
 XX Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGTCATATGCTAAA 21
 DB 30 CCGGGCTGTCATATGCTAAA 50

RESULT 6
 ADQ30647

ID ADQ30647 standard; DNA; 967 BP.

XX ADQ30647;

XX 23-SEP-2004 (first entry)

XX West Nile virus internal diagnosis control sequence.

XX ss; internal control; West Nile Virus; diagnosis.

XX West Nile virus.

XX WO2004055159-A2.

XX 01-JUL-2004.

XX 05-DEC-2003; 2003WO-US038750.

XX 12-DEC-2002; 2002US-0432850P.

XX 20-JUN-2003; 2003US-0480431P.

XX (CHIR) CHIRON CORP.

XX Shyamala V;

XX WPI; 2004-488058/46.

XX New isolated oligonucleotides for accurately diagnosing West Nile virus
 PT infection or for capturing, detecting and quantitating West Nile virus in
 PT blood samples.

XX Claim 27; SEQ ID NO 17; 56pp; English.

XX The invention relates to an isolated oligonucleotide not more than 60
 CC nucleotides in length comprising a nucleotide sequence (S1) of at least
 CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
 CC 20, 21 or 23 bp) given in the specification derived from the West Nile
 CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
 CC identity to the nucleotide sequence of (S1), or complements of (S1) and
 CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
 CC end and/or the 3'-end. The detectable label is a fluorescent label
 CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
 CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
 CC composition and methods are useful for accurately diagnosing West Nile
 CC virus infection or for capturing, detecting and quantitating West Nile
 CC virus in biological samples, particularly blood samples. This sequence
 CC corresponds to an internal control sequence for the detection of WNV
 CC sequences using the oligonucleotides of the invention.

XX Sequence 967 BP; 273 A; 206 C; 272 G; 216 T; 0 U; 0 Other;

XX Query Match 100.0%; Score 21; DB 12; Length 967;

XX Best Local Similarity 100.0%; Pred. No. 0.65;

XX Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGTCATATGCTAAA 21
 DB 129 CCGGGCTGTCATATGCTAAA 149

RESULT 7
 ADR32078

ID ADR32078 standard; DNA; 10945 BP.

XX ADR32078;

XX

DT 18-NOV-2004 (first entry)
 XX Genomic DNA of a West Nile virus.
 XX analysis; target; real time PCR; ds; genomic.
 XX West Nile virus.
 XX WO2004072230-A2.
 XX 26-AUG-2004.
 XX 10-FEB-2004; 2004WO-US002012.
 XX 10-FEB-2003; 2003US-00361004.
 XX (CLEA-) CLEARANT INC.
 XX Mckenney K, Gillmeister L, Marlowe K, Armistead D;
 XX WPI; 2004-625843/60.
 XX Analyzing a target nucleic acid sequence in a biological material by real
 PT time PCR using nucleic acid primers that are separated by at least 750
 PT nucleic acid residues in the target sequence.
 XX Disclosure; SEQ ID NO 5; 96pp; English.
 XX The invention relates to a novel method for analyzing a target nucleic
 CC acid sequence in a biological material. The method comprises adding at
 CC least two nucleic acid primers that hybridise under stringent conditions
 CC to predetermined nucleic acid sequences of the target nucleic acid
 CC sequence that are separated by at least 750 nucleic acid residues,
 CC amplifying the target nucleic acid sequence by PCR, and detecting and
 CC quantifying the target nucleic acid sequence. The methods and
 CC compositions of the present invention are useful for analysing a target
 CC nucleic acid sequence in a biological material by real time PCR using
 CC nucleic acid primers that are separated by at least 750 nucleic acid
 CC residues in the target sequence. This polynucleotide sequence represents
 CC the genomic DNA of a West Nile virus used in the target analysis method
 CC of the invention.
 XX
 SQ Sequence 10945 BP; 2999 A; 2457 C; 3143 G; 2346 T; 0 U; 0 Other;
 Query Match 100.0%; Score 21; DB 13; Length 10945;
 Best Local Similarity 100.0%; Pred. No. 0.93;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 CCGGGCTGTCATATGCTAAA 21
 Db 87 CCGGGCTGTCATATGCTAAA 107
 RESULT 8
 AD67768
 ID AD67768 standard; DNA; 10945 BP.
 XX
 AC AD67768;
 XX
 XX 18-NOV-2004 (first entry)
 DT West Nile virus DNA detected by novel detection method.
 XX ds; detection; pathogen.
 XX
 XX West Nile virus.
 XX WO2004072231-A2.
 XX 26-AUG-2004.
 XX 10-FEB-2004; 2004WO-US002013.
 XX
 XX
 PR 10-FEB-2003; 2003US-00361002.
 XX (CLEA-) CLEARANT INC.
 XX Mckenney K, Gillmeister L, Marlowe K, Armistead D;
 XX WPI; 2004-625844/60.
 XX Determining level of potentially active biological pathogens in
 PT biological material, by adding nucleic acid primer pairs to biological
 PT material, amplifying target nucleic acid by PCR, detecting and
 PT quantifying target nucleic acid.
 XX Disclosure; SEQ ID NO 5; 111pp; English.
 XX The invention relates to a method of determining (MI) level of
 CC potentially active biological pathogens in biological material, involves
 CC adding at least two nucleic acid primer pairs to biological material,
 CC amplifying target nucleic acid sequences by PCR, and detecting and
 CC quantifying target nucleic acid sequences, where quantity of the nucleic
 CC acid sequences is proportional to number of biological pathogens in
 CC biological material. (MI) is useful for determining level of potentially
 CC active biological pathogens in a biological material such as cells,
 CC tissues, blood or blood components, proteins, enzymes, immunoglobulins,
 CC bone marrow, heart valves, cartilage, corneas, nerves, bone, teeth, skin grafts,
 CC lipids, carbohydrates, collagen, chitin and its derivatives, forensic
 CC samples, mummified material, human or animal remains, stem cells, islet
 CC of Langerhans cells, cells for transplantation, red blood cells, white
 CC blood cells or platelets. The biological pathogen is chosen from
 CC bacteria, viruses, fungi and single cell parasites. The biological
 CC pathogen is chosen from Aspergillus, Candida, Histoplasma,
 CC Saccharomyces, Coccidioides, Cryptococcus, Escherichia, Bacillus,
 CC Campylobacter, Helicobacter, Listeria, Clostridium, Streptococcus,
 CC Enterococcus, Staphylococcus, Brucella, Haemophilus, Salmonella,
 CC Yersinia, Pseudomonas, Serratia, Enterobacter, Klebsiella, Proteus,
 CC Citrobacter, Corynebacterium, Propionibacterium and Coxiella. The
 CC biological pathogen is chosen from Adeno-associated virus (AAV),
 CC California encephalitis virus, Coronavirus, Coxsackievirus-A,
 CC Coxsackievirus-B, Eastern equine encephalitis virus (EEEV), Echovirus,
 CC Hantavirus, Hepatitis A virus (HAV), Hepatitis C virus (HCV), Hepatitis
 CC Delta virus (HDV), Hepatitis E virus (HEV), Hepatitis G virus (HGV), HIV,
 CC Human T-lymphotropic virus (HTLV), Influenza virus (Flu virus), Measles
 CC virus (Rubeola), Mumps virus, Norwalk virus, Parainfluenza virus, Polio
 CC virus, Rabies virus, Respiratory Syncytial virus, Rhinovirus, Rubella
 CC virus, Saint Louis encephalitis virus, Western equine encephalitis virus
 CC (WEEV), Yellow fever virus, Adenovirus, Cytomegalovirus (CMV), Epstein-
 CC Barr virus (EBV), Hepatitis B virus (HBV), Herpes simplex virus 1, Herpes
 CC simplex virus 2, Molluscum contagiosum, Papilloma virus (HPV), Smallpox
 CC virus (Variola), Vaccinia virus, Venezuelan equine encephalitis virus
 CC (VEEV), Ebola virus, West Nile virus, Human Parvovirus B19 and Rotavirus.
 CC (MI) is useful for determining the effectiveness of a sterilization
 CC process applied to a biological material. (MI) is useful in determining
 CC whether the biological pathogen is inactive or active. (MI) enables
 CC determination of whether the particular biological pathogen is present in
 CC a biological material as shown by amplification of first target sequence
 CC and whether the biological pathogen is inactive or active. (MI) enables
 CC evaluation of the effectiveness of sterilization processes, and
 CC determination of both the original level and the residual level of
 CC potentially active biological pathogens. This sequence corresponds to a
 CC West Nile virus DNA detected by the method of the invention.
 XX
 SQ Sequence 10945 BP; 2999 A; 2457 C; 3143 G; 2346 T; 0 U; 0 Other;
 Query Match 100.0%; Score 21; DB 13; Length 10945;
 Best Local Similarity 100.0%; Pred. No. 0.93;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 CCGGGCTGTCATATGCTAAA 21
 Db 87 CCGGGCTGTCATATGCTAAA 107

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RESULT 9
ADK13681
ID ADK13681 standard; DNA; 10962 BP.
XX
AC ADK13681;
XX
DT 20-MAY-2004 (first entry)
XX
DE West Nile Virus DNA sequence, SEQ ID 1.
XX
KW Virucide; Immunostimulant; flavivirus;
KW envelope protein domain III polypeptide; envelope protein; gene; ss.
XX
OS West Nile virus.
XX
FH Key Location/Qualifiers
FT CDS 97..10389
FT FT /*tag= a
FT FT /product= "West Nile Virus protein"
XX
PN WO2004016586-A2.
XX
PD 26-FEB-2004.
XX
PF 18-AUG-2003; 2003WO-US025681.
XX
PR 16-AUG-2002; 2002US-0403893P.
PR 06-FEB-2003; 2003US-0445581P.
XX
PA (TEXA ) UNIV TEXAS SYSTEM.
XX
PI Barrett A, Beasley D, Holbrook M;
XX
XX WPI; 2004-203756/19.
DR P-PSDB; ADK13682.
XX
PT Diagnosing flavivirus infection by contacting a sample from a human or
PT animal with a flavivirus envelope protein domain III polypeptide, and
PT detecting formation of an immunocomplex between the envelope protein and
PT antibodies in the sample.
XX
PS Disclosure; SEQ ID NO 1; 110pp; English.
XX
CC The present invention relates to a method for screening for a flavivirus
CC in a subject or animal host. The method comprises: contacting a sample
CC from the subject with a composition comprising a flavivirus envelope
CC protein domain III polypeptide (ADK13683-ADK13701) under conditions that
CC permit formation of specific immunocomplex between an antibody in the
CC sample and the envelope protein domain III polypeptide; and detecting
CC whether a specific immunocomplex is formed. The present sequence is the
CC coding sequence for West Nile Virus protein, from which E protein
CC envelope protein domain III polypeptide (ADK13683) is derived.
XX
SQ Sequence 10962 BP; 2997 A; 2497 C; 3100 G; 2368 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 12; Length 10962;
Best Local Similarity 100.0%; Pred. NO. 0.93;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGTCAATATGCTAAA 21
DB 129 CCGGGCTGTCAATATGCTAAA 149

RESULT 10
ADN98022
ID ADN98022 standard; DNA; 10975 BP.
XX
AC ADN98022;
XX
DT 29-JUL-2004 (first entry)
XX
DE West Nile Virus isolate 2741 complete genome sequence.

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XX
KW ds; West Nile Virus; envelope protein; glycoprotein E; flavivirus;
KW Japanese encephalitis virus; Dengue virus; St Louis encephalitis virus.
XX
OS West Nile virus.
XX
PN WO2004040263-A2.
XX
PD 13-MAY-2004.
XX
PF 31-OCT-2003; 2003WO-US034823.
XX
PR 31-OCT-2002; 2002US-0422755P.
PR 06-JUN-2003; 2003US-0476513P.
XX
XX (HEAL-) HEALTH RES INC.
PA
XX
XX Wong SJ, Pei-Yong S;
XX
XX WPI; 2004-400223/37.
DR GENBANK; AF206518.
XX
XX New diagnostic kit comprising West Nile Virus (WNV) envelope protein
PT reactive with antibody against WNV and cross-reactive with antibody
PT against a flavivirus, useful in diagnosing flavivirus infection caused by
PT DENV, WNV, JEV or SLEV.
XX
XX Disclosure; Fig 37; 212pp; English.
XX
CC The invention relates to a diagnostic kit comprising at least one
CC isolated and purified polypeptide comprising a West Nile Virus (WNV)
CC envelope (E) protein or its immunogenic fragment having a native
CC conformation or non-denatured structure and that is reactive with
CC antibodies against WNV and cross-reactive with antibodies against a
CC flavivirus. The diagnostic kit is useful in diagnosing flavivirus
CC infection caused by DENV, WNV, JEV or SLEV. This sequence corresponds to
CC the complete nucleotide sequence of the WNV isolate 2741.
XX
SQ Sequence 10975 BP; 3007 A; 2460 C; 3149 G; 2359 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 12; Length 10975;
Best Local Similarity 100.0%; Pred. NO. 0.93;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGTCAATATGCTAAA 21
DB 111 CCGGGCTGTCAATATGCTAAA 131

RESULT 11
ABZ68481
ID ABZ68481 standard; DNA; 11029 BP.
XX
AC ABZ68481;
XX
DT 22-APR-2003 (first entry)
XX
DE Nucleotide sequence of the genome of West Nile virus IS-98-ST1.
XX
KW WNV; IS-98-ST1; Flavivirus; infection; encephalitis; gene; ss.
XX
OS West nile virus.
XX
FH Key Location/Qualifiers
FT CDS 97..10397
FT FT /*tag= a
FT FT /product= "polyprotein"
XX
XX WO200281511-A1.
XX
PD 17-OCT-2002.
XX
PF 04-APR-2002; 2002WO-FR001168.

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XX 04-APR-2001; 2001FR-00004599.
PR 06-SEP-2001; 2001FR-00011525.
XX
XX (INSP ) INST PASTEUR.
XX (KIMR-) KIMRON VETERINARY INST.
XX
XX Despres P, Deubel V, Guenet J, Drouet M, Malkinson M, Banet C;
PI Frenkiel M, Courageot M, Coulibaly F, Catteau A, Flamand M, Weber P;
PI Ceccaldi P;
XX
XX WPI; 2003-058498/05.
DR P-PSDB; ABP70647.
DR
XX New neurovirulent strain of West Nile virus, useful in diagnosis and
PT screening for antiviral agents, also related nucleic acids, proteins and
PT antibodies.
XX
XX Claim 1; Page 34-49; 68pp; French.
XX
XX The present sequence represents the genome of a strain of West Nile virus
CC (WNV), designated IS-98-ST1. This strain is a neuroinvasive and
CC neurovirulent strain of WNV. Polynucleotides and polypeptides derived
CC from the IS-98-ST1 genome are useful for diagnosis and prognosis of
CC Flavivirus infection, specifically WNV-mediated encephalitis. They are
CC also useful to raise specific antibodies, for recombinant expression of
CC WNV proteins or peptides (for diagnosis, production of antibodies and
CC identification of specific binding partners in cells), for identifying
CC cellular genes implicated in resistance to viral infection, and for
CC screening for anti-Flavivirus agents
XX
XX Sequence 11029 BP; 3019 A; 2471 C; 3167 G; 2372 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 21; DB 8; Length 11029;
Best Local Similarity 100.0%; Pred. No. 0.93; Length 11029;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CCGGGCTGTCAATATGCTAAA 21
DB 129 CCGGGCTGTCAATATGCTAAA 149
RESULT 12
ABV74821
ID ABV74821 standard; DNA; 11029 BP.
XX
XX AC ABV74821;
XX
XX 28-MAR-2003 (first entry)
XX
XX West Nile virus strain NY99-flamingo 382-99 complete genome.
XX
XX Virucide; hepatotropic; antiinflammatory; antiviral; OAS;
KW 2'-5'-oligoadenylate synthase; Flavivirus infection; gene; ss.
XX
XX West Nile Virus.
XX
XX Key Location/Qualifiers
XX CDS 97..10398
XX /*tag= a
XX /product= "West Nile Virus protein"
XX
XX WO200281741-A2.
XX
XX 17-OCT-2002.
XX
XX 04-APR-2002; 2002WO-FR001169.
XX
XX 04-APR-2001; 2001FR-00004598.
XX
XX (INSP ) INST PASTEUR.
XX (CNRS ) CNRS CENT NAT RECH SCI.
XX

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PI Guenet J, Mashimo T, Simon-Chazottes D, Montagnetelli X;
PI Frenkiel M, Despres P, Deubel V, Bonhomme F, Lucas M;
XX
XX WPI; 2003-058566/05.
DR P-PSDB; ABB98821.
XX
XX Identifying stimulators of oligoadenylate synthase family genes, useful
PT as antiviral agents against Flavivirus, also mutated genes responsible
PT for sensitivity to virus.
XX
XX Example 1; Page 52-67; 93pp; French.
XX
XX The present invention relates to a method for identifying compounds (I)
CC that can stimulate a gene of the OAS (2'-5'-oligoadenylate synthase)
CC family. The method comprises: (a) inducing expression of the OAS gene in
CC a culture of cells from a non-human mammal (Flvr/Flvr or Flvr/Flvr);
CC indicating resistance or sensitivity to Flavivirus infection; (b)
CC treating cells with test compound; and (c) measuring activity of OAS gene
CC relative to a control. (I) are potentially useful as antiviral agents for
CC treating infections by Flaviviruses (e.g. hepatitis C; dengue; yellow
CC fever and various forms of encephalitis). Genomic OAS DNA and derived
CC cDNA, also the encoded proteins, are useful: (a) for treating Flavivirus
CC infection; (b) in screening for anti-Flavivirus agents; and (c) for
CC evaluating sensitivity of subjects to Flavivirus infection and their
CC likely response to interferon treatment, e.g. to identify patients at
CC risk of developing severe forms of such infections. The present sequence
CC is West Nile Virus strain NY99-flamingo 382-99 (IS-98-ST1) complete
CC genome, which was used in an example from the invention. West Nile Virus
CC is one such Flavivirus
XX
XX Sequence 11029 BP; 3019 A; 2471 C; 3167 G; 2372 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 21; DB 10; Length 11029;
Best Local Similarity 100.0%; Pred. No. 0.93; Length 11029;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CCGGGCTGTCAATATGCTAAA 21
DB 129 CCGGGCTGTCAATATGCTAAA 149
RESULT 13
ADN98023
ID ADN98023 standard; DNA; 11029 BP.
XX
XX AC ADN98023;
XX
XX 29-JUL-2004 (first entry)
XX
XX West Nile Virus isolate 3356 complete genome sequence.
XX
XX ds; West Nile Virus; envelope protein; glycoprotein E; flavivirus;
KW Japanese encephalitis virus; Dengue virus; St Louis encephalitis virus.
XX
XX West Nile virus.
XX
XX WO2004040263-A2.
XX
XX 13-MAY-2004.
XX
XX 31-OCT-2003; 2003WO-US034823.
XX
XX 31-OCT-2002; 2002US-0422755P.
XX 06-JUN-2003; 2003US-0476513P.
XX
XX (HEAL-) HEALTH RES INC.
XX
XX Wong SJ, Pei-Yong S;
XX
XX WPI; 2004-400223/37.
XX GENBANK; AF404756.
XX
XX New diagnostic kit comprising West Nile Virus (WNV) envelope protein
PT

```

PT reactive with antibody against WNV and cross-reactive with antibody
 PT against a flavivirus, useful in diagnosing flavivirus infection caused by
 PT DENV, WNV, JEV or SLEV.

XX Disclosure; Fig 38; 212pp; English.

XX The invention relates to a diagnostic kit comprising at least one
 CC isolated and purified polypeptide comprising a West Nile Virus (WNV)
 CC envelope (E) protein or its immunogenic fragment having a native
 CC conformation or non-denatured structure and that is reactive with
 CC antibodies against WNV and cross-reactive with antibodies against a
 CC flavivirus. The diagnostic kit is useful in diagnosing flavivirus
 CC infection caused by DENV, WNV, JEV or SLEV. This sequence corresponds to
 CC the complete nucleotide sequence of the WNV isolate 3356.

XX Sequence 11029 BP; 3017 A; 2466 C; 3172 G; 2374 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 12; Length 11029;
 Best Local Similarity 100.0%; Pred. No. 0.93;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21
 |||||
 Db 129 CCGGGCTGTCATATGCTAAA 149

RESULT 14

ADN36695/c

ID ADN36695 standard; DNA; 21 BP.

XX AC ADN36695;

XX DT 15-JUL-2004 (first entry)

XX West Nile virus detection-related oligonucleotide probe SeqID17.

XX hybridisation assay probe; nucleic acid detection;

XX target-complementary sequence; flavivirus; West Nile virus; WNV;

XX RNA virus; infection; meningitis; encephalitis;

XX high throughput screening; probe; ss.

XX OS West Nile virus.

XX PN WO2004036190-A2.

XX PD 29-APR-2004.

XX PF 10-OCT-2003; 2003WO-US033639.

XX PR 16-OCT-2002; 2002US-0418891P.

XX PR 25-NOV-2002; 2002US-0429006P.

XX PR 24-FEB-2003; 2003US-0449810P.

XX PA (GENP-) GEN-PROBE INC.

XX PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;

XX WPI; 2004-389590/36.

XX New hybridization assay probe comprising target-complementary sequence of
 PT bases, useful in detecting flavivirus, e.g. West Nile virus.

XX Disclosure; SEQ ID NO 17; 135pp; English.

XX This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental

CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening,
 CC the present sequence is that of an oligonucleotide probe which is related
 CC to the invention.

XX Sequence 21 BP; 5 A; 5 C; 4 G; 7 T; 0 U; 0 Other;

Query Match 90.5%; Score 19; DB 12; Length 21;

Best Local Similarity 100.0%; Pred. No. 4.4;

Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 GGGCTGTCATATGCTAAA 21

|||||

Db 21 GGGCTGTCATATGCTAAA 3

RESULT 15

ADN36823/c

ID ADN36823 standard; RNA; 24 BP.

XX AC ADN36823;

XX DT 15-JUL-2004 (first entry)

XX West Nile virus detection-related oligonucleotide probe SeqID145.

XX hybridisation assay probe; nucleic acid detection;

XX target-complementary sequence; flavivirus; West Nile virus; WNV;

XX RNA virus; infection; meningitis; encephalitis;

XX high throughput screening; probe; ss.

XX OS West Nile virus.

XX FH Key Location/Qualifiers

FT modified_base 1..24

FT /*tag= a

FT /mod_base= OTHER

FT /note= "OTHER= 2'-methoxyethoxy (2'-MOE) nucleotides"

XX PN WO2004036190-A2.

XX PD 29-APR-2004.

XX PF 10-OCT-2003; 2003WO-US033639.

XX PR 16-OCT-2002; 2002US-0418891P.

XX PR 25-NOV-2002; 2002US-0429006P.

XX PR 24-FEB-2003; 2003US-0449810P.

XX PA (GENP-) GEN-PROBE INC.

XX PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;

XX WPI; 2004-389590/36.

XX New hybridization assay probe comprising target-complementary sequence of
 PT bases, useful in detecting flavivirus, e.g. West Nile virus.

XX Example 1; SEQ ID NO 145; 135pp; English.

XX This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of an oligonucleotide probe which is related
 CC to the invention.

XX SQ Sequence 24 BP; 5 A; 7 C; 5 G; 0 T; 7 U; 0 Other;
 Query Match 90.5%; Score 19; DB 12; Length 24;
 Best Local Similarity 100.0%; Pred. No. 4.4; Indels 0; Gaps 0;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GGGCTGTCAATATGCTAAA 21
 DB 24 GGGCTGTCAATATGCTAAA 6

RESULT 16
 ADN36707/c
 ID ADN36707 standard; DNA; 48 BP.
 XX AC
 XX ADN36707;
 XX DT 15-JUL-2004 (first entry)
 XX DE West Nile virus detection-related oligonucleotide probe SeqID29.
 XX KW hybridisation assay probe; nucleic acid detection;
 XX KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 XX KW RNA virus; infection; meningitis; encephalitis;
 XX KW high throughput screening; probe; ss.
 XX OS West Nile virus.
 XX OS Enterobacteria phage T7.
 XX FT Key Location/Qualifiers
 FT misc_feature 1..27
 FT /*tag= a
 FT /*note= "T7 promoter sequence"
 FT misc_feature 28..48
 FT /*tag= b
 FT /*note= "WNV-complementary sequence"
 XX FT
 XX PN W02004036190-A2.
 XX PD 29-APR-2004.
 XX PF 10-OCT-2003; 2003WO-US033639.
 XX PR 16-OCT-2002; 2002US-0418891P.
 XX PR 25-NOV-2002; 2002US-0429006P.
 XX PR 24-FEB-2003; 2003US-0449810P.
 XX PA (GENP-) GEN-PROBE INC.
 XX PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
 XX WPI; 2004-389590/36.
 XX PT New hybridization assay probe comprising target-complementary sequence of
 PT bases, useful in detecting flavivirus, e.g. West Nile virus.
 XX PS Disclosure; SEQ ID NO 29; 135pp; English.
 XX CC This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of an oligonucleotide probe which is related
 CC to the invention.

XX SQ Sequence 48 BP; 16 A; 9 C; 9 G; 14 T; 0 U; 0 Other;
 Query Match 90.5%; Score 19; DB 12; Length 48;
 Best Local Similarity 100.0%; Pred. No. 4.9; Indels 0; Gaps 0;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GGGCTGTCAATATGCTAAA 21
 DB 48 GGGCTGTCAATATGCTAAA 30

RESULT 17
 ADN36694/c
 ID ADN36694 standard; DNA; 69 BP.
 XX AC
 XX ADN36694;
 XX DT 15-JUL-2004 (first entry)
 XX DE West Nile virus detection-related oligonucleotide probe SeqID16.
 XX KW hybridisation assay probe; nucleic acid detection;
 XX KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 XX KW RNA virus; infection; meningitis; encephalitis;
 XX KW high throughput screening; probe; ss.
 XX OS West Nile virus.
 XX PN W02004036190-A2.
 XX PD 29-APR-2004.
 XX PF 10-OCT-2003; 2003WO-US033639.
 XX PR 16-OCT-2002; 2002US-0418891P.
 XX PR 25-NOV-2002; 2002US-0429006P.
 XX PR 24-FEB-2003; 2003US-0449810P.
 XX PA (GENP-) GEN-PROBE INC.
 XX PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
 XX WPI; 2004-389590/36.
 XX PT New hybridization assay probe comprising target-complementary sequence of
 PT bases, useful in detecting flavivirus, e.g. West Nile virus.
 XX PS Claim 68; SEQ ID NO 16; 135pp; English.
 XX CC This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of an oligonucleotide probe which is related
 CC to the invention.
 XX SQ Sequence 69 BP; 18 A; 22 C; 14 G; 15 T; 0 U; 0 Other;
 Query Match 90.5%; Score 19; DB 12; Length 69;
 Best Local Similarity 100.0%; Pred. No. 5.2; Indels 0; Gaps 0;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GGGCTGTCAATATGCTAAA 21
 DB 69 GGGCTGTCAATATGCTAAA 51

```

RESULT 18
ADN36696/c
ID   ADN36696 standard; DNA; 20 BP.
XX
XX   AC   ADN36696;
XX
XX   DT   15-JUL-2004 (first entry)
XX
XX   DE   West Nile virus detection-related oligonucleotide probe SeqID18.
XX
XX   KW   hybridisation assay probe; nucleic acid detection;
XX   KW   target-complementary sequence; flavivirus; West Nile virus; WNV;
XX   KW   RNA virus; infection; meningitis; encephalitis;
XX   KW   high throughput screening; probe; ss.
XX
XX   OS   West Nile virus.
XX
XX   PN   WO2004036190-A2.
XX
XX   PD   29-APR-2004.
XX
XX   PF   10-OCT-2003; 2003WO-US033639.
XX
XX   PR   16-OCT-2002; 2002US-0418891P.
XX   PR   25-NOV-2002; 2002US-0429006P.
XX   PR   24-FEB-2003; 2003US-0449810P.
XX
XX   PA   (GENP-) GEN-PROBE INC.
XX
XX   PI   Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX   DR   WPI; 2004-389590/36.
XX
XX   PT   New hybridization assay probe comprising target-complementary sequence of
XX   PT   bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX   PS   Disclosure; SEQ ID NO 18; 135pp; English.
XX
XX   CC   This invention relates to a novel hybridisation assay probe, for
XX   CC   detecting a nucleic acid, which is a probe sequence that comprises a
XX   CC   target-complementary sequence of bases, and optionally one or more base
XX   CC   sequences that are not complementary to the nucleic acid that is to be
XX   CC   detected. The hybridisation assay probes and the kits are useful in
XX   CC   detecting and amplifying a target nucleic acid sequence, for example
XX   CC   flavivirus like West Nile virus, that may be present in a biological
XX   CC   sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX   CC   birds and culex mosquitoes, with humans and horses serving as incidental
XX   CC   hosts. Infection of humans can lead to meningitis or encephalitis. The
XX   CC   invention may allow for accurate and efficient high throughput screening.
XX   CC   The present sequence is that of an oligonucleotide probe which is related
XX   CC   to the invention.
XX
XX   SQ   Sequence 20 BP; 5 A; 4 C; 4 G; 7 T; 0 U; 0 Other;

Query Match      85.7%; Score 18; DB 12; Length 20;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      4 GGCTGTCAATATGCTAAA 21
DB      20 GGCTGTCAATATGCTAAA 3

RESULT 19
ADN36708/c
ID   ADN36708 standard; DNA; 47 BP.
XX
XX   AC   ADN36708;
XX
XX   DT   15-JUL-2004 (first entry)
XX
XX   DE   West Nile virus detection-related oligonucleotide probe SeqID30.

```

```

XX
XX   hybridisation assay probe; nucleic acid detection;
XX   KW   target-complementary sequence; flavivirus; West Nile virus; WNV;
XX   KW   RNA virus; infection; meningitis; encephalitis;
XX   KW   high throughput screening; probe; ss.
XX
XX   OS   West Nile virus.
XX
XX   PN   WO2004036190-A2.
XX
XX   PD   29-APR-2004.
XX
XX   PF   10-OCT-2003; 2003WO-US033639.
XX
XX   PR   16-OCT-2002; 2002US-0418891P.
XX   PR   25-NOV-2002; 2002US-0429006P.
XX   PR   24-FEB-2003; 2003US-0449810P.
XX
XX   PA   (GENP-) GEN-PROBE INC.
XX
XX   PI   Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX   DR   WPI; 2004-389590/36.
XX
XX   PT   New hybridization assay probe comprising target-complementary sequence of
XX   PT   bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX   PS   Disclosure; SEQ ID NO 30; 135pp; English.
XX
XX   CC   This invention relates to a novel hybridisation assay probe, for
XX   CC   detecting a nucleic acid, which is a probe sequence that comprises a
XX   CC   target-complementary sequence of bases, and optionally one or more base
XX   CC   sequences that are not complementary to the nucleic acid that is to be
XX   CC   detected. The hybridisation assay probes and the kits are useful in
XX   CC   detecting and amplifying a target nucleic acid sequence, for example
XX   CC   flavivirus like West Nile virus, that may be present in a biological
XX   CC   sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX   CC   birds and culex mosquitoes, with humans and horses serving as incidental
XX   CC   hosts. Infection of humans can lead to meningitis or encephalitis. The
XX   CC   invention may allow for accurate and efficient high throughput screening.
XX   CC   The present sequence is that of an oligonucleotide probe which is related
XX   CC   to the invention.
XX
XX   SQ   Sequence 47 BP; 16 A; 8 C; 9 G; 14 T; 0 U; 0 Other;

Query Match      85.7%; Score 18; DB 12; Length 47;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      4 GGCTGTCAATATGCTAAA 21
DB      47 GGCTGTCAATATGCTAAA 30

RESULT 20
AAQ22767
ID   AAQ22767 standard; DNA; 4512 BP.
XX
XX   AC   AAQ22767;
XX
XX   DT   12-AUG-1992 (first entry)
XX
XX   DE   JEV Nakayama strain prM, E, NS1, NS2A, NS2B and C coding regions.

```

09-OCT-2003; 2003WO-KR002081.
09-OCT-2002; 2002KR-00061589.
(CIDC-) CID CO LTD.
(LEES/) LEE S H.
Lee SH, Lee Y, Yun S;
WPI; 2004-340933/31.
New Japanese encephalitis virus genomic RNA, useful in developing vaccines for and in diagnosing and treating Japanese encephalitis.
Example 2; Page 145-152; 265pp; English.
The present invention relates to a genomic RNA of the Korean Japanese Encephalitis Virus (JEV) isolate, composed of a 5' and 3' nontranslated region (NTR) and a single polypeptide coding region. The JEV genomic RNA, JEV cDNA and reagents are useful in developing vaccines for and in diagnosing and treating Japanese encephalitis. The present sequence is a sequence of the invention.
Query Match 84.8%; Score 17.8; DB 12; Length 10818;
Best Local Similarity 90.5%; Pred.No. 47;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 1 CCGGGCTGTCATATGCTGAA 21
||||| |||||||
Db 78 CCGGGCTATCAATATGCTGAA 98
RESULT 22
AD007437
ID AD007437 standard; DNA; 10968 BP.
XX
AC AD007437;
XX
DT 15-JUL-2004 (first entry)
XX
DE Japanese Encephalitis virus JEV coding sequence SEQ ID NO: 15.
XX
XX antiinflammatory; neuroprotective; gene therapy;
KW Japanese Encephalitis virus; JEV; ds; gene; vaccine;
KW Japanese encephalitis.
XX
OS Japanese encephalitis virus.
XX
PN WO2004033690-A1.
XX
PD 22-APR-2004.
XX
XX 09-OCT-2003; 2003WO-KR002081.
XX
XX 09-OCT-2002; 2002KR-00061589.
XX
XX (CIDC-) CID CO LTD.
XX (LEES/) LEE S H.
XX
XX Lee SH, Lee Y, Yun S;
XX WPI; 2004-340933/31.
XX
XX New Japanese encephalitis virus genomic RNA, useful in developing vaccines for and in diagnosing and treating Japanese encephalitis.
XX
XX Claim 3; Page 154-161; 265pp; English.
XX
XX The present invention relates to a genomic RNA of the Korean Japanese Encephalitis Virus (JEV) isolate, composed of a 5' and 3' nontranslated region (NTR) and a single polypeptide coding region. The JEV genomic RNA,


```

Qy 1 CCGGGCTCAATAGCTAAA 21
    |||||
Db 128 CCGGCTATCAATAGCTGAA 148

```


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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 17:45:55 ; Search time 1500.84 Seconds
(without alignments)
532.600 Million cell updates/sec

Title: US-10-729-421-34

Perfect score: 21
Sequence: 1 ccgggtgtcaatgctaaa 21

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 100 summaries

Database :

EST: *
1: gb_est1: *
2: gb_est2: *
3: gb_hic: *
4: gb_est3: *
5: gb_est4: *
6: gb_est5: *
7: gb_est6: *
8: gb_gest1: *
9: gb_gest2: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	17.8	84.8	425	7	C0951450 UMC-pd12f
2	17.8	84.8	696	7	CN153647 940784 MA
3	17.8	84.8	696	7	CN155761 943088 MA
4	17.8	84.8	998	9	AG570358 Mus muscu
5	17.4	82.9	429	7	CK699541 ZF101-P00
6	17.4	82.9	693	9	CE434484 tigr-gss-
7	17.4	81.0	1009	9	CNS00HNB Drosophi
8	16.8	80.0	107	9	CC888327 SALK1516
9	16.8	80.0	354	7	CK091023 F039E30.3
10	16.8	80.0	367	7	CK101326 F039P30.5
11	16.8	80.0	407	4	BG037611 dc53e09.Y
12	16.8	80.0	501	7	CN489600 Mdfw20191
13	16.8	80.0	516	8	AZ981379 2M0258113
14	16.8	80.0	562	5	B0815985 N058F04 P
15	16.8	80.0	620	6	CA821839 RSH08C06
16	16.8	80.0	623	8	AZ832056 2M0112109
17	16.8	80.0	631	7	CK317796 B9P01h01
18	16.8	80.0	635	5	B0863468 S028D11 P
19	16.8	80.0	751	7	CV257160 WS0245.B2
20	16.8	80.0	790	9	CR268537 Reverse s
21	16.8	80.0	900	9	CR016265 Forward s
22	16.8	80.0	986	5	BQ220271 AGENCOURT
23	16.4	78.1	298	9	CE199293 tigr-gss-
24	16.4	78.1	363	1	AL926049 AL926049

1	502	1	AI512957	AI512957 LD45125.5
2	522	5	BX723363	BX723363 BX723363
3	604	8	BH039924	BH039924 RPCI-24-2
4	686	7	CO125121	CO125121 GR_EB08G
5	704	9	CC870522	CC870522 NDL_123C7
6	815	5	BU381267	BU381267 603861468
7	851	5	BX723398	BX723398 BX723398
8	851	5	BX723398	BX723398 saf21f09.9
9	223	4	BI320339	BI320339 k8k0712.f
10	403	7	CF190539	CF190539 Mdfw20230
11	405	7	CN496900	CN496900 Mdfw20230
12	432	7	CO285402	CO285402 EK168918.
13	503	1	AA918584	AA918584 OL51a09.8
14	537	8	AZ713066	AZ713066 RPCI-24-6
15	543	6	BY566461	BY566461 BY566461
16	551	8	AQ425764	AQ425764 CITBI-E1-
17	560	9	CE166745	CE166745 tigr-gss-
18	570	5	BX711144	BX711144 BX711144
19	573	7	CN955009	CN955009 Mdfw2050m
20	573	7	BF260707	BF260707 HVSMEF002
21	578	4	BG000437	BG000437 MR3-GN015
22	592	4	BM440929	BM440929 EBto08 SQ
23	601	2	BF614355	BF614355 de05e10.Y
24	618	2	BF676682	BF676682 602086323
25	621	6	CA694474	CA694474 wlmk4_pk0
26	637	7	CN910418	CN910418 030128ABL
27	651	4	BJ589589	BJ589589 BJ589589
28	652	7	CF365321	CF365321 835940 MA
29	659	4	BJ591475	BJ591475 BJ591475
30	698	4	BJ589606	BJ589606 BJ589606
31	721	9	AG417300	AG417300 Mus muscu
32	732	9	AG558847	AG558847 Mus muscu
33	743	9	AG548062	AG548062 Mus muscu
34	744	9	AG566555	AG566555 Mus muscu
35	756	4	BJ594699	BJ594699 BJ594699
36	757	7	CO802419	CO802419 AGENCOURT
37	800	9	BX997985	BX997985 Reverse B
38	824	8	AZ187754	AZ187754 SP_1009.B
39	859	8	BZ701219	BZ701219 PUCOC20TD
40	860	8	BZ704489	BZ704489 PUCCN20TD
41	863	7	CF374332	CF374332 AGENCOURT
42	886	5	BX851663	BX851663 BX851663
43	932	9	CL240184	CL240184 ZMMBB0508
44	936	5	BQ688620	BQ688620 AGENCOURT
45	939	9	CNS04CON	AL283856 Tetraodon
46	942	6	CD245530	CD245530 AGENCOURT
47	953	9	CG094980	CG094980 PUKBV26TB
48	1009	9	CG094979	CG094979 PUKBV26TB
49	1016	9	CL997901	CL997901 ZMMBB0019
50	1049	9	CL391702	CL391702 ZMMBB0019
51	1058	8	CC221736	CC221736 CH261-921
52	1062	8	CC221316	CC221316 CH261-83N
53	391	4	BI028233	BI028233 CM4-WT028
54	402	6	CB218410	CB218410 NISC nb08
55	535	8	AQ226974	AQ226974 HS_2013.A
56	588	9	CE471568	CE471568 tigr-gss-
57	657	8	BH352121	BH352121 CH230-220
58	774	7	CK448276	CK448276 pncs900ad
59	929	9	CNS041CB	AL292196 Tetraodon
60	959	9	CL477040	CL477040 SAIL 266
61	1050	9	CL046976	CL046976 CH216-65F
62	220	6	CD112634	CD112634 ME1-0024T
63	234	7	R40815	R40815 yf82b01.s1
64	246	7	R61488	R61488 yf815g09.s1
65	268	5	BP751208	BP751208 BP751208
66	270	1	AV210948	AV210948 AV210948
67	273	7	H09942	H09942 ym01c11.s1
68	286	1	AA958331	AA958331 MAAD0514.
69	294	1	AV228707	AV228707 AV228707
70	295	8	CC072824	CC072824 CSU-K33r.
71	306	4	BI126863	BI126863 I081P66P
72	328	4	BI126556	BI126556 I077P15P
73	340	8	AZ783049	AZ783049 2M0024J20
74	350	1	AU233672	AU233672 AU233672

98 15.8 75.2 355 1 AU278073 AU278073
 C 99 15.8 75.2 388 9 CE627437 tigr-gss-
 c 100 15.8 75.2 430 8 AQ437513 HS_5133_B

ALIGNMENTS

RESULT 1
 C0951450
 LOCUS
 DEFINITION UMC-pd12fol-008-a09 Day 12 ovarian follicle pd12fol Sus scrofa cDNA
 3', mRNA sequence.

ACCESSION C0951450
 VERSION C0951450.1 GI:51328453
 KEYWORDS EST.
 SOURCE Sus scrofa (pig)

ORGANISM

Sus scrofa
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.

REFERENCE

1 (bases 1 to 425)
 Jiang, H., Whitworth, K.M., Bivens, N.J., Ries, J.E., Woods, R.J.,
 Forrester, L.J., Springer, G.K., Mathialagan, N., Agca, C.,
 Prather, R.S. and Lucy, M.C.

AUTHORS

Large-scale Generation and Analysis of Expressed Sequence Tags from
 Porcine Ovary

TITLE

Unpublished (2004)

JOURNAL

Contact: DNA Core Facility (Swine Project)
 Animal Science - RS Prather
 University of Missouri-Columbia

COMMENT

M616 Medical Sciences Bldg., Columbia, MO 65212, USA
 Tel: (573)882-0428
 Fax: (573)884-5552
 Email: porcine@net.missouri.edu
 POLYA=NO.

FEATURES

source

1..425
 Location/Qualifiers
 /organism="Sus scrofa"
 /mol_type="mRNA"
 /db_xref="taxon:9823"
 /dev_stage="Day 12 ovarian follicle"
 /clone_lib="pd12fol"
 /note="Vector: T3T7PAC; Funding: A grant from the Monsanto
 Company to the University of Missouri. Genetic Source:
 Ovarian tissue (whole ovary, dissected follicles, or
 corpora lutea) was collected from crossbred pigs (Sus
 scrofa domestica), frozen in liquid nitrogen shortly after
 collection, and stored at -80 degrees Celsius until RNA
 extraction. The tissue from several individual pigs was
 pooled for the purpose of RNA extraction. The specific
 tissues collected were fetal whole ovary; neonatal whole
 ovary; prepubertal whole ovary; 2, 4, 6 and 8 mm growing
 follicles; Day 0 follicles; Day 5 small antral follicles
 and corpora lutea; Day 12 corpora lutea and Day 12
 follicles. More information regarding the methods can be
 found at:
 http://genome.mnet.missouri.edu/Swine/Methods.html.
 Library Construction (Standard Protocol): All procedures
 discussed in this section have been described in detail
 elsewhere (Soares et al., 1994; Bonaldo et al., 1996;
 Jiang et al., 2001). Total cellular RNA from each sample
 was isolated by using STAT-60 reagent (Tel-test,
 Friendswood, TX) and the poly(A)+ RNA was obtained by two
 rounds of purification with the Oligotex mRNA isolation
 kit (Qiagen) according to the manufacturer's instructions.
 The libraries were constructed essentially as described by
 the manufacturer's instructions provided with the
 Superscript Plasmid System (Invitrogen, cat. no.
 18248-013). Briefly, 1mg of poly(A)+ RNA will be annealed
 at 37 degrees Celsius with 10mg of NotI-tag-drl8
 oligonucleotide (GTCGTCGGCGGC-tag-T18) and reverse
 transcribed at 37 degrees Celsius with Superscript II
 (Invitrogen) reverse transcriptase (Jiang et al., 2001).

The 'tag' represents a tissue/stage-specific ten-base
 sequence identifier
 (http://genome.uiowa.edu/pubsoft/software.html) present in
 the oligonucleotide used to prime first-strand synthesis.
 Second strand synthesis was performed with T4 DNA
 polymerase in the presence of DNA ligase and RNase H.
 After second strand synthesis, the double-stranded cDNAs
 was ligated to SalI adapters (Invitrogen-Life
 Technologies) and digested with NotI. The cDNAs will be
 size selected by passage through cDNA size fractionation
 columns (Invitrogen-Life Technologies). The cDNAs derived
 from each developmental stage of a particular tissue were
 mixed on an equimolar basis and ligated directionally into
 the NotI and SalI sites of the pSPORT1 vector
 (Invitrogen). After ligation of the inserts, the plasmids
 will be electroporated into DH10B bacteria. Preliminary
 Library Characterization: Randomly chosen clones from each
 library were analyzed by restriction digestion to
 determine average insert size (96 clones) and by
 sequencing (~4 96-well plates) to confirm library quality
 [e.g. the presence of short polyA+ tails, genomic DNA
 contamination (must be <1%), ribosomal RNA clones (must be
 <1%), etc.] and to provide a sequence database
 representing the predominant clones in each library. The
 clones were sequenced at the University of
 Missouri-Columbia DNA Core Facility. Bioinformatics work
 was performed by GK Springer's bioinformatics group (WG
 Spollen, JE Ries, A Guillen, AA Khambati, RV Patel, CM
 Topinka, SB Bhuiyan) in Computer Science and Health
 Management and Informatics Departments at the University
 of Missouri-Columbia. Clone Requests: Requests for clones
 should be made to the Director of the University of
 Missouri DNA Core facility at: porcine@net.missouri.edu.
 Citations: 1. Bonaldo MF, Lennon G, Soares MB.
 Normalization and Subtraction: Two approaches to
 facilitate gene discovery. Genome Res, 1996; 6:791-806.
 2. Jiang H, Bivens NJ, Ries JE, Whitworth KM, Green JA,
 Forrester LJ, Springer GK, Didion BA, Mathialagan N,
 Prather RS, Lucy MC (2001) Constructing cDNA libraries
 with fewer clones that contain long poly(cA) tails.
 Biotechniques 31:38-42. 3. Soares MB, MF Bonaldo, P
 Jelenne, L Su, L Lawton, A Efstratiadis. 1994.
 Construction and characterization of a normalized cDNA
 library. Proc Natl Acad Sci, 91:9228-9232.
 TAG_TISSUE=Day 12 ovarian follicle
 TAG_SEQ=Not found"

ORIGIN

Query Match 84.8%; Score 17.8; DB 7; Length 425;
 Best Local Similarity 90.5%; Pred. No. 2.2e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CCGGGCTGTCATATGCTAAA 21

|||||||
 Db 49 CCGGGCTGCGCATATGCTAAA 69

RESULT 2
 CNI53647/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Sus scrofa

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.

1 (bases 1 to 696)

Smith, T.P.L., Freking, B.A., Ford, J.J., Vallet, J.L., Wise, T.A.,

Nonneman, D.J., Wray, J.E. and Keele, J.W.

Porcine EST collection using a normalized library constructed from

embryos representing early developmental stages

CNI53647 696 bp mRNA linear EST 02-APR-2004
 940784 MARC 4PIG Sus scrofa cDNA 3', mRNA sequence.

CNI53647

CNI53647.1 GI:46169077

EST.

Sus scrofa (pig)

Sus scrofa

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.

1 (bases 1 to 696)

Smith, T.P.L., Freking, B.A., Ford, J.J., Vallet, J.L., Wise, T.A.,

Nonneman, D.J., Wray, J.E. and Keele, J.W.

Porcine EST collection using a normalized library constructed from

embryos representing early developmental stages

JOURNAL COMMENT
Unpublished (2003)
Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called with phred v0.020425.c and trimmed with the aid of the trim_alt option. Vector identified with cross_match v0.990329.
Plate: TWM8048 row: D column: 8
Seq primer: TAGAAGGCACATGTCGAGG.
Location/Qualifiers
1..696
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/tissue_type="pooled"
/lab_host="DH10B"
/clone_lib="WARC 4PIG"
/note="vector: pCDNA3.1; Site 1: EcoRI; Site 2: NotI; Library made with combined RNA from day-10, day-13, day-15, day-25, and day-30 whole embryos."

FEATURES
source
Query Match 84.8%; Score 17.8; DB 7; Length 696;
Best Local Similarity 90.5%; Pred. No. 2.3e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

ORIGIN
Query Match 84.8%; Score 17.8; DB 7; Length 696;
Best Local Similarity 90.5%; Pred. No. 2.3e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CCGGGCTGCAATATGCTTAA 21
|||||
Db 488 CCGGGCTGCGGATATGCTTAA 468
|||||

RESULT 3
LOCUS CN155761 696 bp mRNA linear EST 02-APR-2004
DEFINITION 943088 WARC 4PIG Sus scrofa cDNA 5', mRNA sequence.
ACCESSION CN155761
VERSION 1 GI:46170191
KEYWORDS EST.
SOURCE Sus scrofa (pig)
ORGANISM Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE 1 (bases 1 to 696)
AUTHORS Smith,T.P.L., Freking,B.A., Ford,J.J., Vallet,J.L., Wise,T.A., Nonneman,D.J., Wray,J.E. and Keele,J.W.
TITLE Porcine EST collection using a normalized library constructed from embryos representing early developmental stages
JOURNAL Unpublished (2003)
COMMENT Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called with phred v0.020425.c and trimmed with the aid of the trim_alt option. Vector identified with cross_match v0.990329.
Plate: TWM8048 row: D column: 8
Seq primer: GTAATACGACTCACTATAGG.
Location/Qualifiers
1..696
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/tissue_type="pooled"
/lab_host="DH10B"
/clone_lib="WARC 4PIG"
/note="vector: pCDNA3.1; Site 1: EcoRI; Site 2: NotI; Library made with combined RNA from day-10, day-13, day-15, day-25, and day-30 whole embryos."

JOURNAL COMMENT
Unpublished (2003)
Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called with phred v0.020425.c and trimmed with the aid of the trim_alt option. Vector identified with cross_match v0.990329.
Plate: TWM8048 row: D column: 8
Seq primer: TAGAAGGCACATGTCGAGG.
Location/Qualifiers
1..696
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/tissue_type="pooled"
/lab_host="DH10B"
/clone_lib="WARC 4PIG"
/note="vector: pCDNA3.1; Site 1: EcoRI; Site 2: NotI; Library made with combined RNA from day-10, day-13, day-15, day-25, and day-30 whole embryos."

FEATURES
source
Query Match 84.8%; Score 17.8; DB 7; Length 696;
Best Local Similarity 90.5%; Pred. No. 2.3e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

ORIGIN
Query Match 84.8%; Score 17.8; DB 7; Length 696;
Best Local Similarity 90.5%; Pred. No. 2.3e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CCGGGCTGCAATATGCTTAA 21
|||||
Db 209 CCGGGCTGCGGATATGCTTAA 229
|||||

RESULT 4
LOCUS AG570358 998 bp DNA linear GSS 05-JUN-2004
DEFINITION Mus musculus molossinus DNA, clone:MSMg01-492K10.TJ, genomic survey sequence.
ACCESSION AG570358
VERSION 1 GI:48331078
KEYWORDS GSS.
SOURCE Mus musculus molossinus
ORGANISM Mus musculus molossinus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1
AUTHORS Hattori,M., Toyoda,A., Noguchi,H., Kojima,T. and Sakaki,Y.
TITLE BAC end Sequences of Library MSMg01
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 998)
AUTHORS Hattori,M., Toyoda,A., Noguchi,H., Kojima,T. and Sakaki,Y.
TITLE Direct Submission
JOURNAL Submitted (17-NOV-2003) Masahira Hattori, The Institute of Physical and Chemical Research (RIKEN), Genomic Sciences Center (GSC); 1-7-22 Suehiro-chou,Tsukumi-ku, Yokohama, Kanagawa 230-0045, Japan [E-mail:hattori@gsc.riken.jp, URL:http://hgp.gsc.riken.go.jp/, Tel:81-45-503-9111, Fax:81-45-503-9170]
COMMENT Clones are derived from the mouse BAC library MSMg01. For BAC library availability, please contact Kuniya Abe (abe@tc.riken.jp). Tsukuba Institute, Bio Resource Center, The Institute of Physical and Chemical Research (RIKEN) 3-1-1 Koyadai, Tsukuba, 305-0074 Japan
phone: 81-298-36-9189, fax: 81-298-36-9199
e-mail: abe@tc.riken.jp
PRIMERS
Sequencing : TJ
LIBRARY : pBACE3.6
Vector : pBACE3.6
R.Site 1 : EcoRI
R.Site 2 : EcoRI.
Location/Qualifiers
1..998
/organism="Mus musculus molossinus"
/mol_type="genomic DNA"
/sub_species="molossinus"
/db_xref="taxon:57486"
/clone="MSMg01-492K10.TJ"
/sex="male"
/tissue_type="mixture of kidney and spleen"
/clone_lib="MSMg01 Mouse Male BAC Library"

FEATURES
source
Query Match 84.8%; Score 17.8; DB 9; Length 998;
Best Local Similarity 90.5%; Pred. No. 2.4e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

ORIGIN
Query Match 84.8%; Score 17.8; DB 9; Length 998;
Best Local Similarity 90.5%; Pred. No. 2.4e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CCGGGCTGCAATATGCTTAA 21
|||||
Db 193 CCGGGCTGCAATATGCTTAA 213
|||||

RESULT 5
LOCUS CK699541 429 bp mRNA linear EST 30-MAR-2004
DEFINITION ZF101-P00082-DEPE-F_G24 GISZF001_ra Danio rerio cDNA clone

```

IMAGE:7165298 5', mRNA sequence.
ACCESSION CK699541
VERSION CK699541.1 GI:42451877
KEYWORDS EST.
SOURCE Danio rerio (zebrafish)
ORGANISM Danio rerio
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
TITLE Cypriniformes; Cyprinidae; Danio.
JOURNAL 1 (bases 1 to 429)
COMMENT Wei, C., Mathavan, S., Thoreau, H., Lim, L., Lee, C. and Ruan, Y.
Genome Institute of Singapore, Zebrafish Gene Collection
Unpublished (2004)
Contact: Ruan Y
Cloning and Sequencing
Genome Institute of Singapore
60 Biopolis Street, #02-01, Genome, Singapore 138672
Tel: +65 6478 8073
Fax: +65 6478 9059
Email: ruanyj@gis.a-star.edu.sg
GIS Clone ID: ZF101-P00082-BR2_G24
PCR Primers
FORWARD: M13
BACKWARD: M13
Plate: ZF101-P00082-BR2 row: G column: 24
Seq primer: CCGCATACTGTATAGCA
High quality sequence stop: 429.
FEATURES
source
1. 429
Location/Qualifiers
/organism="Danio rerio"
/mol_type="mRNA"
/strain="Singapore local strain"
/db_xref="taxon:7955"
/clone="IMAGE:7165298"
/tissue_type="Embryo"
/dev_stage="7 Different embryonic Stages(From just
fertilized Embryos to 72 hours just hatched baby fish)"
/lab_host="DH108"
/clone_lib="GISZP001.ra"
/notes="Vector: pDNR-LIB; Site 1: Sfi A (GCCATTACGGCC);
Site 2: Sfi B (GCCGAGCGGCC); Priming method: Sfi-(dT)30
Primed ; Priming sequence:
5.ATTCTAGCGCGGAGCGCGGCACATG(T)30VN ; Directionally
cloned, 5' cloning site: Sfi A site GCCATTACGGCC ; 5'
linker/adaptor sequence: 5.AAGCAGTGTATCAACGACGATGGCC ;
3' cloning site: Sfi B site GCCGAGCGGCC ; 3'
linker/adaptor sequence: same as the priming sequence ;
Average insert size: 2kb ; For PCR insert analysis: Use
M13 Forward and reverse primers ; Library Amplified ;
Recombinants (inserts): 98% ; Library complexity: 5x106 ;
Full-length construction (method): SMART, a Clontech
method The pooled tissue RNA was collected and used to
construct full length enriched cDNA library and also
served as template to synthesize complex first strand cDNA
probe. Two high density colony arrays were made from over
110K cDNA clones and hybridized with the probes. Low
intensity clones were selected as they represented rare
expressed clones. The hybridization intensities for all
clones span from 0 to 1.8 million counts and the low
abundant class ranged from 0 to 13,000."
ORIGIN
Query Match 82.9%; Score 17.4; DB 7; Length 429;
Best Local Similarity 94.7%; Pred. No. 3.6e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 3 GGGCTGTCATATGCTAAA 21
Db 3 GGGCTGTCATATGCTAAA 21
RESULT 6
LOCUS CNS000HNB
DEFINITION Drosophila melanogaster genome survey sequence T7 end of BAC:
BAC35008 of RPCI-98 library from Drosophila melanogaster (fruit
fly), genomic survey sequence.
ACCESSION AL073822
VERSION AL073822.1 GI:4953796
KEYWORDS GSS.
SOURCE Drosophila melanogaster (fruit fly)
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 1009)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
- Web : www.genoscope.cns.fr)
COMMENT Determination of this BAC-end sequence was carried out as part of a
collaboration with the Berkeley Drosophila Genome Project (BDGP).
The BDGP is constructing a physical map of the Drosophila
melanogaster genome using these BACs. For further information
please see http://www.fruitfly.org The BDGP Drosophila
melanogaster BAC library was prepared by Kazutoyo Osoegawa and
Aaron Mammoser in Pieter de Jong's laboratory in the Department of

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LOCUS CE434484
DEFINITION tigr-gss-dog-17000363221477 Dog Library Canis familiaris genomic,
Genomic survey sequence.
ACCESSION CE434484
VERSION CE434484.1 GI:36711024
KEYWORDS GSS.
SOURCE Canis familiaris (dog)
ORGANISM Canis familiaris
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
REFERENCE 1 (bases 1 to 693)
AUTHORS Kirkness, E.F., Bafna, V., Halpern, A.L., Levy, S., Remington, K.,
Rusch, D.B., Delcher, A.L., Pop, M., Wang, W., Fraser, C.M. and
Venter, J.C.
TITLE The dog genome: survey sequencing and comparative analysis
JOURNAL Science 301 (5641), 1898-1903 (2003)
MEDLINE 22875432
PUBMED 14512627
COMMENT Contact: Kirkness EF
The Institute for Genomic Research
Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive,
Rockville, MD 20850, USA
Tel: 301-838-0200
Fax: 301-838-0208
Email: ekirknes@tigr.org
Class: shotgun.
Location/Qualifiers
1. 693
/organism="Canis familiaris"
/mol_type="genomic DNA"
/strain="Standard Poodle"
/db_xref="taxon:9615"
/clone_lib="dog Library"
/notes="Site 1: BstXI; Libraries were prepared from
peripheral blood"
ORIGIN
Query Match 82.9%; Score 17.4; DB 9; Length 693;
Best Local Similarity 94.7%; Pred. No. 3.8e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 3 GGGCTGTCATATGCTAAA 21
Db 326 GGGCTGTCATATGCTAAA 308
RESULT 7
LOCUS CNS000HNB
DEFINITION Drosophila melanogaster genome survey sequence T7 end of BAC:
BAC35008 of RPCI-98 library from Drosophila melanogaster (fruit
fly), genomic survey sequence.
ACCESSION AL073822
VERSION AL073822.1 GI:4953796
KEYWORDS GSS.
SOURCE Drosophila melanogaster (fruit fly)
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 1009)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
- Web : www.genoscope.cns.fr)
COMMENT Determination of this BAC-end sequence was carried out as part of a
collaboration with the Berkeley Drosophila Genome Project (BDGP).
The BDGP is constructing a physical map of the Drosophila
melanogaster genome using these BACs. For further information
please see http://www.fruitfly.org The BDGP Drosophila
melanogaster BAC library was prepared by Kazutoyo Osoegawa and
Aaron Mammoser in Pieter de Jong's laboratory in the Department of

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Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain Y2; cn bw sp, the same strain used for the BDGP's p1 and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

FEATURES

source

1..1009
Location/Qualifiers
/organism="Drosophila melanogaster"
/mol_type="genomic DNA"
/db_xref="taxon:7227"
/clone="BACR35D08"
/clone_lib="RPCI-98"
/note="end : T7"

ORIGIN

Query Match 81.0%; Score 17; DB 9; Length 1009;
Best Local Similarity 85.7%; Pred. No. 6.4e+02;
Matches 18; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21

Db 133 CCGGCTGTCAATATACHAA 153

RESULT 8

CC888327/c

LOCUS 107 bp DNA linear GSS 31-JUL-2003
DEFINITION SALK_151698.23.35-x Arabidopsis thaliana TDNA insertion lines
Arabidopsis thaliana genomic clone SALK_151698.23.35.x, genomic survey sequence.

ACCESSION CC888327.1 GI:33364847

VERSION

KEYWORDS

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM

Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.

1 (bases 1 to 107)

Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R.,
Gadrinab,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L.,
Shinn,P., Zimmerman,J. and Ecker,J.R.

A Sequence-Indexed Library of Insertion Mutations in the

Arabidopsis Genome

Unpublished (2001)

Contact: Joseph R. Ecker

Salk Institute Genomic Analysis Laboratory (SIGNAL)

The Salk Institute for Biological Studies

10010 N. Torrey Pines Road, La Jolla, CA 92037, USA

Tel: 858 453 4100 x1752

Fax: 858 558 6379

Email: ecker@salk.edu

This is single pass sequence recovered from the left border of

TDNA.

Class: TDNA tagged.

FEATURES

source

Location/Qualifiers
1..107
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/ecotype="Col-0"
/db_xref="taxon:3702"
/clone="SALK_151698.23.35.x"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
notes="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"

ORIGIN

Query Match 80.0%; Score 16.8; DB 9; Length 107;
Best Local Similarity 90.0%; Pred. No. 6.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 20

Db 79 CCGGGCTGTCATATGCTAAA 60

RESULT 9

CK091023

LOCUS

DEFINITION F039P30.3Pr Populus flower cDNA library Populus balsamifera subsp.
trichocarpa cDNA clone F039P30 3', mRNA sequence.

CK091023

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Populus balsamifera subsp. trichocarpa (Populus trichocarpa)
Populus balsamifera subsp. trichocarpa
Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids I; Malpighiales; Salicaceae; Populus.

REFERENCE

AUTHORS

1 (bases 1 to 354)
Sterky, F., Bhalerai, R.R., Unneberg, P., Segerman, B., Nilsson, P.,
Brunner, A.M., Campaa, L., Jonsson-Lindvall, J., Tandré, K.,
Strauss, S.H., Sundberg, B., Gustafsson, P., Uhlen, M., Bhalerai, R.P.,
Nilsson, O., Sandberg, G., Karlsson, J., Lundberg, J. and Jansson, S.

A Populus EST resource for functional genomics

Unpublished (2003)

Other ESTs: F039P30Y, F039P30.5Pr

Contact: Bo Segerman

Umea Plant Science Center, Department of Plant Physiology

Umea University

901 87 Umea, Sweden

Tel: +46 90 786 5279

Fax: +46 90 786 6676

Email: bo.segerman@plantphys.umu.se.

Location/Qualifiers

1..354

/organism="Populus balsamifera subsp. trichocarpa"

/mol_type="mRNA"

/sub_species="trichocarpa"

/db_xref="taxon:3694"

/clone="F039P30"

/tissue_type="floral buds"

/clone_lib="Populus flower cDNA library"

/note="Organ: flower"

ORIGIN

Query Match 80.0%; Score 16.8; DB 7; Length 354;
Best Local Similarity 90.0%; Pred. No. 7.3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 CCGGGCTGTCATATGCTAAA 21

Db 213 CCGGGCTTCAAAATGCTAAA 232

RESULT 10

CK101326/c

LOCUS

DEFINITION F039P30.5Pr Populus flower cDNA library Populus balsamifera subsp.
trichocarpa cDNA clone F039P30 5', mRNA sequence.

CK101326

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Populus balsamifera subsp. trichocarpa (Populus trichocarpa)

Populus balsamifera subsp. trichocarpa

Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids I; Malpighiales; Salicaceae; Populus.

1 (bases 1 to 367)

pollination) ACCGA18(T) insert. Double stranded cDNAs were size selected (more than 450 bp), adapted with EcoRI adapters at both ends and then digested with NotI. The cDNAs were then directionally cloned into EcoRI-NotI digested pBS II SK(+) phagemid vector (Stratagene). Identification of adaptors and tags in 5'-end sequenced clones: <Vector>...TAGCTT<End Vector><Start EcoRI adaptor>GATATCGAATTCATGTGTGGG <End EcoRI adaptor><Start Insert>...AAAAAAAAAAAAAAAA<End NotI site>Vector>GGCGCGCCACCGCGG... The total number of white colony forming units (cfu) in the primary library before amplification was 1.1x10⁶ cfu (colony forming units). The background of empty clones was less than 1%. Purified plasmid DNA from the primary library was converted to single-stranded circles and used as a template for PCR amplification using the T7 and T3 priming sites flanking the cloned cDNA inserts. The purified PCR products, representing the entire cloned cDNA population, were used as a driver for normalization. Hybridization between the single-stranded library and the PCR products was carried out for 44 hours at 30C. Unhybridized single-stranded DNA circles were separated from hybridized DNA rendered partially double-stranded and electroporated into DH10B cells to generate the normalized library. The total number of clones with insert was 9x10⁶ cfu. Background of empty clones was less than 1%.

ORIGIN

Query Match 80.0%; Score 16.8; DB 7; Length 501;
Best Local Similarity 85.7%; Pred. No. 7.6e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CCGGCTGCTCAATATGCTAAA 21
|||||
Db 332 CCGGCTGCTCCATTGCTGAAA 312

RESULT 13

AZ981379/c
LOCUS 516 bp DNA linear GSS 27-APR-2001
DEFINITION 2M0258113R Mouse 10kb plasmid UUGC2M library Mus musculus genomic clone UUGC2M0258113 R, genomic survey sequence.

ACCESSION AZ981379
VERSION GI:13852606
KEYWORDS GSS.
SOURCE Mus musculus (house mouse)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 516)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D.,Weiss,R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

JOURNAL

COMMENT Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177

Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0258 row: 1 column: 13
Seq primer: CACACGGAACAGCTATGACC
Class: plasmid ends
High quality sequence stop: 516.
Location/Qualifiers

FEATURES

source

1. 516
/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC2M0258113"
/sex="female"
/lab_host="E. coli strain XL10-Gold, Tl-resistant, P-"
/clone_lib="Mouse 10kb plasmid UUGC2M library"
/notes="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (female) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (gi|4732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

ORIGIN

Query Match 80.0%; Score 16.8; DB 8; Length 516;
Best Local Similarity 90.0%; Pred. No. 7.6e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 CCGGCTGCTCAATATGCTAAA 21
|||||
Db 30 CTGGCTGCTCAATATGCTACA 11

RESULT 14

BUB15985/c
LOCUS 562 bp mRNA linear EST 15-OCT-2002
DEFINITION N058F04 Populus bark cDNA library Populus tremula x Populus tremuloides cDNA 5 prime, mRNA sequence.

ACCESSION BUB15985
VERSION GI:23975607
KEYWORDS EST.

SOURCE

Populus tremula x Populus tremuloides
Populus tremula x Populus tremuloides
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosids; eurosids I; Malpighiales; Salicaceae; Populus. 1 (bases 1 to 562)

REFERENCE

Unneberg,P., Bhalerao,R.R., Jansson,S. and Sterky,P.
The poplar tree transcriptome: Analysis of expressed sequence tags from multiple libraries
Unpublished (2002)

JOURNAL

COMMENT Contact: BHALERAO RUPALI R.
Umea Plant Science Center
Department of Plant Physiology
University of Umea, 901 87 Umea, Sweden
Tel: +46 90 786 5279
Fax: +46 90 786 6676
Email: rupali.bhalerao@plantphys.umu.se.
Location/Qualifiers

FEATURES

1. 562
/organism="Populus tremula x Populus tremuloides"
/mol_type="mRNA"
/db_xref="taxon:47664"
/tissue_type="bark"
/clone_lib="Populus bark cDNA library"

ORIGIN

Query Match	80.0%;	Score 16.8;	DB 5;	Length 562;	
Best Local Similarity	90.0%;	Pred. No. 7.7e+02;			
Matches 18;	Conservative 0;	Mismatches 2;	Indels 0;	Gaps 0;	
QY	2	CGGGCTGTCAATATGCTAAA 21			
Db	293	CGGGCTTCAAGATGCTAAA 274			
RESULT 15					
CA821839/c					
LOCUS					
DEFINITION					
CA821839					
DESCRIPTION					
CA821839					
ACCESSION					
VERSION					
KEYWORDS					
SOURCE					
ORGANISM					
REFERENCE					
AUTHORS					
TITLE					
JOURNAL					
COMMENT					
FEATURES					
source					
1..620					
/organism="Populus balsamifera subsp. trichocarpa x					
Populus deltoides"					
/mol_type="mRNA"					
/cultiivar="Beaupre"					
/db_xref="taxon:3695"					
/dev_stage="two-month-old"					
/clone_lib="two-month-old roots from clone 'Beaupre' grown					
for 19 days under restricted irrigation"					
/notes="Organ: root; Vector: pTriplex2; cDNA library of					
roots from two-month-old Populus trichocarpa Torr. & Gray					
x deltoides Bartr. Ex Marshall (clone 'Beaupre') grown for					
19 days under restricted irrigation to reach 50% of the					
transpiration rate of fully watered plants. The cDNA					
library was constructed from 1 ug of total RNA using the					
SMART cDNA synthesis kit (Clontech, Palo Alto, CA, USA)					
according to the manufacturer's instructions. The					
resulting cDNA was packed into lambda phages using the					
Gigapack III Gold packaging kit (Stratagene, La Jolla,					
CA). The pTriplex2 phagemid clones in Escherichia coli					
were obtained by using the mass in vivo excision protocol					
according to the manufacturer's instructions (Clontech)."					
ORIGIN					
Query Match	80.0%;	Score 16.8;	DB 6;	Length 620;	
Best Local Similarity	90.0%;	Pred. No. 7.8e+02;			
Matches 18;	Conservative 0;	Mismatches 2;	Indels 0;	Gaps 0;	
QY	2	CGGGCTGTCAATATGCTAAA 21			
Db	389	CGGGCTTCAAGATGCTAAA 370			
RESULT 16					
AZ832056					
LOCUS					
DEFINITION					
AZ832056					
ACCESSION					
VERSION					
KEYWORDS					
SOURCE					
ORGANISM					
REFERENCE					
AUTHORS					
TITLE					
JOURNAL					
COMMENT					
FEATURES					
source					
1..623					
/organism="Mus musculus"					
/mol_type="genomic DNA"					
/strain="C57BL/6J"					
/db_xref="taxon:10090"					
/clone="UUGC2M0112L09"					
/sex="Male"					
/lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"					
/clone_lib="Mouse 10kb plasmid UUGC1M library"					
/note="Vector: PWB42nv; Purified genomic DNA from M.					
musculus C57BL/6J (male) was obtained from the Jackson					
Laboratory Mouse DNA Resource					
(http://www.jax.org/resources/documents/dnares/). The DNA					
was hydrodynamically sheared by repeated passage through a					
0.005 inch orifice at constant velocity. The sheared DNA					
was blunt end-repaired with T4 DNA polymerase and T4					
polynucleotide kinase. Adaptor oligonucleotides were					
ligated to the blunt ends in high molar excess. The					
adaptor DNA was purified and size-selected for a 9.5 to					
10.5 kb range using preparative agarose gel					
electrophoresis. Vector DNA was prepared from a derivative					
of pWD42 (gi 4732114 gb AF129072.1), a copy-number					
inducible derivative of plasmid R1. The vector was ligated					
with adaptors complementary to the insert adaptors and					
purified. The sheared, adaptor mouse DNA was annealed to					
adaptor vector DNA, and transformed into					
chemically-competent E. coli XL10-Gold (Stratagene) cells					
and selected for ampicillin resistance."					
ORIGIN					
Query Match	80.0%;	Score 16.8;	DB 8;	Length 623;	
Best Local Similarity	90.0%;	Pred. No. 7.8e+02;			
Matches 18;	Conservative 0;	Mismatches 2;	Indels 0;	Gaps 0;	
QY	2	CGGGCTGTCAATATGCTAAA 21			
Db	563	CTGGCTGTCAATATGCTACA 582			
RESULT 17					
CK317796/c					

LOCUS
DEFINITION B9P01h01 Populus stem seasonal library Populus deltoides cDNA, mRNA
ACCESSION CK317796 631 bp mRNA linear EST 11-MAY-2004
VERSION CK317796
KEYWORDS CK317796.1 GI:47106219
SOURCE EST.
ORGANISM Populus deltoides
 Populus deltoides
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 rosids; eurosids I; Malpighiales; Salicaceae; Populus.
REFERENCE 1 (bases 1 to 631)
AUTHORS Park, S. and Han, K.-H.
TITLE Gene expression profile during seasonal growth cycle in poplar tree
JOURNAL Unpublished (2003)
COMMENT Contact: Kyung-Hwan Han
 Department of Forestry
 Michigan State University
 126 Natural Resources, East Lansing, MI 48824-1222, USA
 Tel: 517 353 4751
 Fax: 517 432 1143
 Email: hanky@msu.edu.

FEATURES
 source
 Location/Qualifiers
 1..631
 /organism="Populus deltoides"
 /mol_type="mRNA"
 /strain="ILL-129"
 /db_xref="taxon:3696"
 /tissue_type="stem"
 /dev_stage="1 year old"
 /clone_lib="Populus stem seasonal library"

ORIGIN
 Query Match 80.0%; Score 16.8; DB 7; Length 631;
 Best Local Similarity 90.0%; Pred. No. 7.8e+02;
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 2 CGGGCTGTCGAATATGCTAAA 21
 ||||| ||||| ||||| ||||| |||||
 Db 355 CGGGCTTCAAGATGCTAAA 336

RESULT 18
LOCUS BU863468/c
DEFINITION S028D11 Populus imbibed seed cDNA library Populus tremula cDNA 5
 prime, mRNA sequence.
ACCESSION BU863468 635 bp mRNA linear EST 16-OCT-2002
VERSION BU863468.1 GI:24049528
KEYWORDS EST.
SOURCE Populus tremula
 Populus tremula
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 rosids; eurosids I; Malpighiales; Salicaceae; Populus.
REFERENCE 1 (bases 1 to 635)
AUTHORS Unneberg, P., Bhalerao, R.R., Jansson, S. and Sterky, P.
TITLE The poplar tree transcriptome: Analysis of expressed sequence tags
 from multiple libraries
JOURNAL Unpublished (2002)
COMMENT Contact: BHALERAO RUPALI R.
 Umea Plant Science Center
 Department of Plant Physiology
 University of Umea, 901 87 Umea, Sweden
 Tel: +46 90 786 5279
 Fax: +46 90 786 6676
 Email: rupali.bhalerao@plantphys.umu.se.

FEATURES
 source
 Location/Qualifiers
 1..635
 /organism="Populus tremula"
 /mol_type="mRNA"
 /db_xref="taxon:113636"
 /tissue_type="imbibed seed"

LOCUS
DEFINITION /clone_lib="Populus imbibed seed cDNA library"
ACCESSION 80.0%; Score 16.8; DB 5; Length 635;
VERSION 90.0%; Pred. No. 7.8e+02;
KEYWORDS Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
SOURCE 2 CGGGCTGTCGAATATGCTAAA 21
 ||||| ||||| ||||| ||||| |||||
ORGANISM Db 340 CGGGCTTCAAGATGCTAAA 321

RESULT 19
LOCUS CV257160 751 bp mRNA linear EST 22-SEP-2004
DEFINITION WS0245_B21_N14 PTxD-ICC-N-A-14 Populus balsamifera subsp.
 trichocarpa x Populus deltoides cDNA clone WS0245_N14 3', mRNA
 sequence.
ACCESSION CV257160
VERSION CV257160.1 GI:52510135
KEYWORDS EST.
SOURCE Populus balsamifera subsp. trichocarpa x Populus deltoides
 Populus balsamifera subsp. trichocarpa x Populus deltoides
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 rosids; eurosids I; Malpighiales; Salicaceae; Populus.
REFERENCE 1 (bases 1 to 751)
AUTHORS Ralph, S., Cooper, D., Kolosova, N., Oddy, C., Butterfield, Y.,
 Kirkpatrick, R., Liu, J., Palmquist, D., Stott, J., Barber, S., Yang, G.,
 Babakaliff, R., Brown-John, M., Chand, S., Featherstone, R., Mason, A.,
 Mayo, M., Moran, J., Olson, T., Wong, D., Ritland, C.E., Siddiqui, A.,
 Holt, R., Jones, S., Marra, M., Ellis, B.E., Douglas, C., Ritland, K. and
 Bohlmann, J.
TITLE The poplar transcriptome: Analysis of expressed sequence tags from
 multiple cDNA libraries
JOURNAL Unpublished (2004)
COMMENT Contact: Joerg Bohlmann
 Genome BC forest genomics program
 University of British Columbia
 UBC Biotechnology Laboratory, 6174 University Boulevard, Rm. 237,
 Vancouver, British Columbia, Canada, V6T 1Z3
 Tel: 1-604-822-0282
 Fax: 1-604-822-6097
 Email: bohlmann@interchange.ubc.ca
 Plate: WS0245 row: N column: 14
 High quality sequence stop: 751
 POLYA=Yes.

FEATURES
 source
 Location/Qualifiers
 1..751
 /organism="Populus balsamifera subsp. trichocarpa x
 Populus deltoides"
 /mol_type="mRNA"
 /cultivar="Hil-11"
 /db_xref="taxon:3695"
 /clone="WS0245_N14"
 /sex="Male"
 /lab_host="E. coli DH10B T1 phage resistant cells"
 /clone_lib="PTxD-ICC-N-A-14"
 /note="Vector: pBluescript II SK (+) XR; Site_1: EcoRI (5'
 end of cDNA); Site_2: XhoI (3' end of cDNA); Cultured
 cells [de Sa MM et al. (1992) Plant Physiology 98:728-737]
 were grown in media (45mL) supplemented with either 50uM
 salicylic acid, 50uM benzothiadiazole, 50uM methyl
 jasmonate, 20ug chitosan or 200uL of Pollacia radiosa
 extract. Cells were harvested after a 3 hour treatment,
 along with untreated control cells. mRNA was isolated from
 each tissue source independently and equal quantities of
 mRNA from each tissue were then pooled. cDNA was prepared
 from 5 micrograms of mRNA and directionally ligated into
 the pBluescript II SK (+) XR vector using the pBluescript
 II XR cDNA Library Construction Kit according to
 manufacturer's instructions with modifications
 (Stratagene). Plasmid DNA was then transformed by

electroporation into DH10B cells (Invitrogen) for propagation. Normalization was applied according to published methods [Donaldo M.F. et al. (1996) Genome Research 6(9):791] in order to reduce the abundance of highly expressed transcripts."

ORIGIN

Query Match 80.0%; Score 16.8; DB 7; Length 751;
Best Local Similarity 90.0%; Pred. No. 8e+02; Indels 0; Gaps 0;
Matches 18; Conservative 0; Mismatches 2;

QY 2 CCGGCTGTCAATATGCTAAA 21
||||| ||||| ||||| |||||

Db 491 CCGGCTTCAGATGCTAAA 510
||||| ||||| ||||| |||||

RESULT 20

CR268537/c

LOCUS CR268537 790 bp DNA linear GSS 06-JUL-2004
DEFINITION Reverse strand read from insert in 5'HPRT insertion targeting and chromosome engineering clone MHPN344a20, genomic survey sequence.

CR268537

ACCESSION CR268537.1 GI:50047390

VERSION GSS; genome survey sequence; MICER.

KEYWORDS Mus musculus (house mouse)

SOURCE

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 790)

AUTHORS Adams,D.J., Biggs,P.J., Cox,A.V., Davies,R.M., van der Weyden,L.,

Jonkers,J., Smith,J., Plumb,R.W., Taylor,R.G., Nishijima,I., Yu,Y.,

Rogers,J., and Bradley,A.

Direct Submission

TITLE Submitted (20-FEB-2004) Sanger Centre, Hinxton, Cambridgeshire,

JOURNAL CB10 1SA, UK. http://www.sanger.ac.uk/MICER

FEATURES

source

1..790

/organism="Mus musculus"

/mol_type="genomic DNA"

/db_xref="taxon:10090"

/clone="MHPN344a20"

/clone_lib="MHPN"

ORIGIN

Query Match 80.0%; Score 16.8; DB 9; Length 790;
Best Local Similarity 90.0%; Pred. No. 8e+02; Indels 0; Gaps 0;
Matches 18; Conservative 0; Mismatches 2;

QY 2 CCGGCTGTCAATATGCTAAA 21
||||| ||||| ||||| |||||

Db 361 CTGGCTGTCAATATGCTACA 342
||||| ||||| ||||| |||||

RESULT 21

CR016265

LOCUS CR016265 900 bp DNA linear GSS 05-JUL-2004

DEFINITION Forward strand read from insert in 3'HPRT insertion targeting and chromosome engineering clone MHP8d17, genomic survey sequence.

CR016265

ACCESSION CR016265.1 GI:49749320

VERSION GSS; genome survey sequence; MICER.

KEYWORDS Mus musculus (house mouse)

SOURCE

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 900)

AUTHORS Adams,D.J., Biggs,P.J., Cox,A.V., Davies,R.M., van der Weyden,L.,

Jonkers,J., Smith,J., Plumb,R.W., Taylor,R.G., Nishijima,I., Yu,Y.,

Rogers,J., and Bradley,A.

Direct Submission

TITLE Submitted (20-FEB-2004) Sanger Centre, Hinxton, Cambridgeshire,

JOURNAL CB10 1SA, UK. http://www.sanger.ac.uk/MICER

FEATURES

Location/Qualifiers

source

1..900
/organism="Mus musculus"
/mol_type="genomic DNA"
/db_xref="taxon:10090"
/clone="MHP8d17"
/clone_lib="MHP8"

ORIGIN

Query Match 80.0%; Score 16.8; DB 9; Length 900;
Best Local Similarity 90.0%; Pred. No. 8.1e+02; Indels 0; Gaps 0;
Matches 18; Conservative 0; Mismatches 2;

QY 1 CCGGGCTGTCAATATGCTAA 20
||||| ||||| ||||| |||||

Db 145 CCGGGCTGTCAAGATGCTCA 164
||||| ||||| ||||| |||||

RESULT 22

BQ220271/c

LOCUS BQ220271 986 bp mRNA linear EST 02-MAY-2002

DEFINITION AGENCOURT_7572589 NIH_MGC_92 Homo sapiens cDNA clone IMAGE:6044520

5', mRNA sequence.

ACCESSION BQ220271

VERSION BQ220271.1 GI:20401671

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 986)

AUTHORS NIH-MGC http://mgi.nci.nih.gov/.

National Institutes of Health, Mammalian Gene Collection (MGC)

Unpublished (1999)

JOURNAL

COMMENT

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

Tissue Procurement: ATCC

cDNA Library Preparation: Life Technologies, Inc.

DNA Sequencing by: The I.M.A.G.E. Consortium (LLNL)

Clone distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

http://image.llnl.gov

Plate: LLAM13287 row: e column: 01

High quality sequence stop: 149.

FEATURES

source

1..986

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:6044520"

/tissue_type="embryonal carcinoma, cell line"

/lab_host="DH10B (phage-resistant)"

/clone_lib="NIH_MGC_92"

/note="Organ: testis; Vector: pCMV-SPORT6; Site 1: NotI;

Site 2: SalI; Cloned unidirectionally; oligo-dT primed.

Average insert size 2.5 kb. Library enriched for

full-length clones and constructed by Life Technologies.

Note: this is a NIH_MGC Library."

ORIGIN

Query Match 80.0%; Score 16.8; DB 5; Length 986;
Best Local Similarity 90.0%; Pred. No. 8.2e+02; Indels 0; Gaps 0;
Matches 18; Conservative 0; Mismatches 2;

QY 2 CCGGGCTGTCAATATGCTAAA 21
||||| ||||| ||||| |||||

Db 758 CAGGCTGTGATGCTAAA 739
||||| ||||| ||||| |||||

RESULT 23

CE199293

LOCUS CE199293 298 bp DNA linear GSS 25-SEP-2003

DEFINITION tigr-gss-dog-17000372211318 Dog Library Canis familiaris genomic,

```

ACCESSION      CE199293
VERSION        CE199293.1 GI:35354946
KEYWORDS      GSS.
SOURCE        Canis familiaris (dog)
ORGANISM      Canis familiaris
               Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
               Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
REFERENCE      1 (bases 1 to 298)
AUTHORS      Kirkness, E.F., Bafna, V., Halpern, A.L., Levy, S., Remington, K.,
               Rusch, D.B., Delcher, A.L., Pop, M., Wang, W., Fraser, C.M. and
               Venter, J.C.
TITLE        The dog genome: survey sequencing and comparative analysis
JOURNAL        Science 301 (5641), 1898-1903 (2003)
MEDLINE       22875432
PUBMED        14512627
COMMENT      Contact: Kirkness EF
               The Institute for Genomic Research
               Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive,
               Rockville, MD 20850, USA
               Tel: 301-838-0200
               Fax: 301-838-0208
               Email: ekirknes@tigr.org
               Class: shotgun.
FEATURES      Location/Qualifiers
               source
               1..298
               /organism="Canis familiaris"
               /mol_type="genomic DNA"
               /strain="Standard Poodle"
               /db_xref="taxon:9615"
               /clone_lib="Dog Library"
               /notes="Site 1: BstXI; Libraries were prepared from
               peripheral blood"
ORIGIN
Query Match      78.1%; Score 16.4; DB 9; Length 298;
Best Local Similarity 94.4%; Pred. No. 1.2e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      4 GGCTGTCATATGCTAAA 21
        |||||
Db      241 GGCTGTAAATATGCTAAA 258

RESULT 24
AL926049/c      363 bp mRNA linear EST 06-JUL-2004
LOCUS          AL926049 PUR-Z1+22 Danio rerio cDNA clone 164-D08-2, mRNA sequence.
DEFINITION     AL926049
ACCESSION      AL926049
VERSION        AL926049.1 GI:23192629
KEYWORDS      EST.
SOURCE        Danio rerio (zebrafish)
ORGANISM      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
               Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
               Cypriniformes; Cyprinidae; Danio.
REFERENCE      1 (bases 1 to 363)
AUTHORS      Lo, J., Lee, S., Xu, M., Liu, F., Ruan, H., Eun, A., He, Y., Ma, W.,
               Wang, W., Wen, Z. and Peng, J.
TITLE        15000 unique zebrafish EST clusters and their future use in
               microarray for profiling gene expression patterns during
               embryogenesis
JOURNAL        Genome Res. 13 (3), 455-466 (2003)
MEDLINE       22505427
PUBMED        12618376
COMMENT      Contact: Peng J
               Lab of Functional Genomics
               Institute of Molecular and Cell Biology
               30 Medical Drive, Singapore, 117609, Singapore
               Email: pengjr@mcb.a-star.edu.sg
               Clone requests: info@openbiosystems.com
               Open Biosystems,
               6705 Odyssey Drive, Huntsville, AL 35806.

```

```

FEATURES      Location/Qualifiers
               source
               1..363
               /organism="Danio rerio"
               /mol_type="mRNA"
               /strain="local wildtype"
               /db_xref="taxon:7955"
               /clone="164-D08-2"
               /tissue_type="whole embryo or fish"
               /dev_stage="mixed stages"
               /clone_lib="PUR-Z1+22"
ORIGIN
Query Match      78.1%; Score 16.4; DB 1; Length 363;
Best Local Similarity 94.4%; Pred. No. 1.2e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      4 GGCTGTCATATGCTAAA 21
        |||||
Db      239 GGCTGTCAGATGCTAAA 222

RESULT 25
AL512957/c      502 bp mRNA linear EST 19-APR-2001
LOCUS          LD45125.Sprime LD Drosophila melanogaster embryo pOT2 Drosophila
DEFINITION     melanogaster cDNA clone LD45125 Sprime similar to D83486: Su(fu)
               PBgn0005355 PID:gi208417 SPTREMBL:Q27279, mRNA sequence.
ACCESSION      AL512957
VERSION        AL512957.1 GI:4422375
KEYWORDS      EST.
SOURCE        Drosophila melanogaster (fruit fly)
ORGANISM      Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
               Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
               Ephydroidea; Drosophilidae; Drosophila.
REFERENCE      1 (bases 1 to 502)
AUTHORS      Harvey, D., Brokstein, P., Hong, L., Evans-Holm, M., Su, C., Teang, G.,
               Lewis, S. and Rubin, G.M.
TITLE        BDGP/HMI Drosophila EST Project
JOURNAL        Unpublished (2001)
COMMENT      Contact: Stapleton, M.
               BDGP
               Lawrence Berkeley National Lab
               One Cyclotron Rd, Berkeley, CA 94720, USA
               Fax: 510 486 6798
               Email: http://www.fruitfly.org/EST, est@fruitfly.berkeley.edu
               Plate: 451 row: C column: 1
               High quality sequence stop: 179.
FEATURES      Location/Qualifiers
               source
               1..502
               /organism="Drosophila melanogaster"
               /mol_type="mRNA"
               /db_xref="taxon:7227"
               /clone="LD45125"
               /sex="male and female"
               /dev_stage="0 to 24 hours mixed stage embryonic"
               /lab_host="XL1 Blue"
               /clone_lib="LD Drosophila melanogaster embryo pOT2"
               /note="Organ: embryo; Vector: pOT2; Site 1: EcoRI; Site 2:
               XhoI; Sized fractionated cDNAs were directly ligated into
               pOT2."
ORIGIN
Query Match      78.1%; Score 16.4; DB 1; Length 502;
Best Local Similarity 94.4%; Pred. No. 1.2e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 CCGGGCTGTCATATGCT 18
        |||||
Db      363 CCGGGTTGTCATATGCT 346

```

Search completed: September 6, 2005, 21:55:48

Job time : 1507.84 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 21:56:10 ; Search time 1636 Seconds
(without alignments)
535.133 Million cell updates/sec

Title: US-10-729-421-8
Perfect score: 23
Sequence: 1 tcatgactgaattccggtcttt 23

Scoring table: OLIGO NUC
Gapop_60.0 , Gapext 60.0

Searched: 34239544 seqs, 19032134700 residues
Word size : 10

Total number of hits satisfying chosen parameters: 61

Minimum DB seq length: 0
Maximum DB seq length: 60

Post-processing: Listing first 45 summaries

Database : EST:*
1: gb_est1:*
2: gb_est2:*
3: gb_hic:*
4: gb_est3:*
5: gb_est4:*
6: gb_est5:*
7: gb_est6:*
8: gb_gss1:*
9: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
C 1	12	52.2	55	1	AA619888	AA619888 vl58h07.8
C 2	11	47.8	29	8	CC456749	CC456749 SALK_1002
C 3	11	47.8	33	8	AZ817376	AZ817376 2M0086N22
C 4	11	47.8	41	8	BH791854	BH791854 SALK_0618
C 5	11	47.8	41	8	BH812487	BH812487 SALK_0618
C 6	11	47.8	50	4	BG153713	BG153713 nag57g06.
C 7	11	47.8	52	2	AW249907	AW249907 2821659.3
C 8	11	47.8	55	7	CN564600	CN564600 tag20b04.
C 9	11	47.8	57	8	BH918919	BH918919 3526_1_63
C 10	10	43.5	19	9	AJ588850	AJ588850 Arabidops
C 11	10	43.5	25	9	AG197078	AG197078 Pan trogl
C 12	10	43.5	34	9	BX660142	BX660142 Arabidops
C 13	10	43.5	36	8	BH810737	BH810737 SALK_0511
C 14	10	43.5	36	9	DM8546528	AJ564528 Drosophil
C 15	10	43.5	37	1	AA972482	AA972482 op42d03.s
C 16	10	43.5	37	1	AI937582	AI937582 wp81b11.x
C 17	10	43.5	37	7	U44311	U44311 ENU44311 As
C 18	10	43.5	37	9	BX531847	BX531847 Arabidops
C 19	10	43.5	39	1	AV833097	AV833097 Arabidops
C 20	10	43.5	39	5	BX568339	BX568339 EX568339
C 21	10	43.5	39	9	TA160H03P	AL473258 T. brucei
C 22	10	43.5	41	9	CR397281	CR397281 Arabidops
C 23	10	43.5	42	9	BX943895	BX943895 Arabidops
C 24	10	43.5	44	9	AL752437	AL752437 Arabidops

C 25	10	43.5	46	6	CF049474	CF049474 QCL37a04.
C 26	10	43.5	46	7	CN753215	CN753215 APHL3LD-X
C 27	10	43.5	47	8	BH000511	BH000511 2M0288M20
C 28	10	43.5	48	4	BG253356	BG253356 602362952
C 29	10	43.5	48	9	BX945507	BX945507 Arabidops
C 30	10	43.5	48	9	CR356946	CR356946 Arabidops
C 31	10	43.5	48	9	CL528770	CL528770 ASV9G01.F
C 32	10	43.5	49	5	BQ100687	BQ100687 1722G04.X
C 33	10	43.5	49	6	CB305243	CB305243 3'EST-Nf1
C 34	10	43.5	49	7	CO733304	CO733304 SILT02C05
C 35	10	43.5	50	1	AU103081	AU103081 AU103081
C 36	10	43.5	50	1	AU103082	AU103082 AU103082
C 37	10	43.5	50	8	BH612727	BH612727 SALK_0331
C 38	10	43.5	52	1	AA068274	AA068274 mm53C01.X
C 39	10	43.5	52	2	BF632337	BF632337 NF018503D
C 40	10	43.5	52	8	AZ629385	AZ629385 1M0482N16
C 41	10	43.5	52	9	BX122966	BX122966 Danio rer
C 42	10	43.5	53	4	BG524434	BG524434 42-53 Ste
C 43	10	43.5	53	7	CN870218	CN870218 001204AAO
C 44	10	43.5	53	8	BH252021	BH252021 SALK_0124
C 45	10	43.5	54	8	AZ576149	AZ576149 AST-T11C0

ALIGNMENTS

RESULT 1
AA619888/c
LOCUS
DEFINITION
v158h07.s1 Knowles Solter mouse 2 cell Mus musculus cDNA clone
IMAGE: 976477 5', mRNA sequence.
ACCESSION
AA619888
VERSION
AA619888.1 GI:2523764
KEYWORDS
EST.
SOURCE
Mus musculus (house mouse)
ORGANISM
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
1 (bases 1 to 55)
AUTHORS
Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T., Geisels, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M., Schellenger, K., Steptoe, M., Tan, F., Underwood, K., Moore, B., Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and Waterston, R.
TITLE
The WashU-HMI Mouse EST Project
JOURNAL
Unpublished (1996)
COMMENT
Contact: Marra M/Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@watson.wustl.edu
This clone is available royalty-free through LILN; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
MGI:557205

FEATURES
source

Location/Qualifiers
1..55
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J x DBA/2J F1"
/db_xref="taxon:10090"
/clone="IMAGE:976477"
/tissue_type="embryo"
/dev_stages="2-cell"
/lab_host="DH10B"
/clone_lib="Knowles Solter mouse 2 cell"
/note="Organ: embryo; Vector: pBluescribe (modified); Site: 1: MluI; Site 2: SalI; Cloned unidirectionally from mRNA prepared from 13,500 2-cell stage embryos. Primer: SalI (dr): 5'-CGGTGACGCGACCGTGTGTGT-3'. CDNA5 were cloned into the MluI/SalI sites of a modified pBluescribe vector using commercial linkers (NEB)."


```

ORIGIN
  Average insert size: 1.2 kb."
  Query Match      52.2%; Score 12; DB 1; Length 55;
  Best Local Similarity 100.0%; Pred. No. 7.5e+03;
  Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GACTGCAATTC 16
   |||||
Db 21 GACTGCAATTC 10

RESULT 2
CC456749          29 bp DNA linear GSS 30-MAY-2003
LOCUS SALK_100255.25.15.x Arabidopsis thaliana TDNA insertion lines
DEFINITION Arabidopsis thaliana genomic clone SALK_100255.25.15.x, genomic
survey sequence.
ACCESSION CC456749
VERSION CC456749
KEYWORDS CC456749.1 GI:31217770
SOURCE CC456749
ORGANISM Arabidopsis thaliana (thale cress)
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 29)
Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R.,
Gadzinab,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L.,
Shinn,P., Zimmerman,J. and Ecker,J.R.
A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
Unpublished (2001)
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGNAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 433 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of
TDNA. This sequence lies within 300 bases of the 5' end of
At2g27775.
Class: TDNA tagged.
FEATURES
  Location/Qualifiers
    1..29
      /organism="Arabidopsis thaliana"
      /mol_type="genomic DNA"
      /scotye="Col-0"
      /db_xref="taxon:3702"
      /clone="SALK_100255.25.15.x"
      /clone_lib="Arabidopsis thaliana TDNA insertion lines"
      /notes="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"

ORIGIN
  Query Match      47.8%; Score 11; DB 8; Length 29;
  Best Local Similarity 100.0%; Pred. No. 3.1e+04;
  Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 9 GCAATTCGGGT 19
   |||||
Db 12 GCAATTCGGGT 22

RESULT 3
AZ817376          33 bp DNA linear GSS 20-FEB-2001
LOCUS SALK_061833.40.05.x Arabidopsis thaliana TDNA insertion lines
DEFINITION

clone UUGC2M0086N22 R, genomic survey sequence.
AZ817376          GI:12987380
GSS.
Mus musculus (house mouse)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 33)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T.,
Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von
Niederhausern,A. and Wright,D., Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0086 row: N column: 22
Seq primer: CACACAGGAACACAGCATGACC
Class: plasmid ends
High quality sequence stop: 33.
Location/Qualifiers
  1..33
    /organism="Mus musculus"
    /mol_type="genomic DNA"
    /strain="C57BL/6J"
    /db_xref="taxon:10090"
    /clone="UUGC2M0086N22"
    /sex="Male"
    /lab_host="E. Coli strain XL10-Gold, Tl-resistant, P-"
    /clone_lib="Mouse 10kb plasmid UUGC1M library"
    /note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 [gi|4732114|gb|AF129072.1], a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

ORIGIN
  Query Match      47.8%; Score 11; DB 8; Length 33;
  Best Local Similarity 100.0%; Pred. No. 3.1e+04;
  Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 TGACTGCAATT 14
   |||||
Db 23 TGACTGCAATT 33

RESULT 4
BH791854/c        41 bp DNA linear GSS 02-APR-2002
LOCUS SALK_061833.40.05.x Arabidopsis thaliana TDNA insertion lines
DEFINITION

```

Arabidopsis thaliana genomic clone SALK_061833.40.05.x, genomic survey sequence.

ACCESSION BH791854
 VERSION BH791854.1 GI:19886147
 KEYWORDS GSS.
 SOURCE Arabidopsis thaliana (thale cress)
 ORGANISM Arabidopsis thaliana

REFERENCE 1 (bases 1 to 41)
 AUTHORS Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P., Zimmerman,J. and Ecker,J.R.
 TITLE A Sequence-Indexed Library of Insertion Mutations in the Arabidopsis Genome
 JOURNAL Unpublished (2001)
 COMMENT Contact: Joseph R. Ecker
 The Salk Institute Genomic Analysis Laboratory (SIGNAL)
 10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
 Tel: 858 453 4100 x1752
 Fax: 858 558 6379
 Email: ecker@salk.edu
 This is single pass sequence recovered from the left border of TDNA.

Class: TDNA tagged.
 Location/Qualifiers
 1..41
 /organism="Arabidopsis thaliana"
 /mol_type="genomic DNA"
 /ecotype="Col-0"
 /db_xref="taxon:3702"
 /clone="SALK_061833"
 /clone_lib="Arabidopsis thaliana TDNA insertion lines"
 /note="PCR was performed on Arabidopsis thaliana lines each of which contains one or more TDNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at http://signal.salk.edu/tdna_protocols.html"

ORIGIN
 Query Match 47.8%; Score 11; DB 8; Length 41;
 Best Local Similarity 100.0%; Pred. No. 3.1e+04;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 ATGACTGCAAT 13
 |||||
 Db 38 ATGACTGCAAT 28

RESULT 5
 BH812487/c
 LOCUS
 DEFINITION 41 bp DNA linear GSS 02-MAY-2002
 SALK_061833 Arabidopsis thaliana TDNA insertion lines Arabidopsis thaliana genomic clone SALK_061833, genomic survey sequence.
 BH812487
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
 Arabidopsis thaliana (thale cress)
 Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 1 (bases 1 to 41)
 Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P., Zimmerman,J. and Ecker,J.R.
 TITLE A Sequence-Indexed Library of Insertion Mutations in the Arabidopsis Genome
 JOURNAL Unpublished (2001)
 COMMENT Contact: Joseph R. Ecker

Salk Institute Genomic Analysis Laboratory (SIGNAL)
 The Salk Institute for Biological Studies
 10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
 Tel: 858 453 4100 x1752
 Fax: 858 558 6379
 Email: ecker@salk.edu
 This is single pass sequence recovered from the left border of TDNA.

Class: TDNA tagged.
 Location/Qualifiers
 1..41
 /organism="Arabidopsis thaliana"
 /mol_type="genomic DNA"
 /ecotype="Col-0"
 /db_xref="taxon:3702"
 /clone="SALK_061833"
 /clone_lib="Arabidopsis thaliana TDNA insertion lines"
 /note="PCR was performed on Arabidopsis thaliana lines each of which contains one or more TDNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at http://signal.salk.edu/tdna_protocols.html"

ORIGIN
 Query Match 47.8%; Score 11; DB 8; Length 41;
 Best Local Similarity 100.0%; Pred. No. 3.1e+04;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 ATGACTGCAAT 13
 |||||
 Db 38 ATGACTGCAAT 28

RESULT 6
 BG153713/c
 LOCUS
 DEFINITION 50 bp mRNA linear EST 05-FEB-2001
 nag57906.x1 NCI_CGAP_Co26 Homo sapiens cDNA clone IMAGE:4225738 3', mRNA sequence.
 BG153713
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
 Homo sapiens (human)
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 50)
 NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
 Unpublished (1997)
 Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 cDNA Library Preparation: David B. Krizman, Ph.D.
 DNA Sequencing by: The I.M.A.G.E. Consortium/LLNL
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL, send email to: info@image.llnl.gov
 Seq primer: -40UP from Gibco.
 Location/Qualifiers
 1..50
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:4225738"
 /tissue_type="normal colonic mucosa"
 /lab_host="DH10B"
 /clone_lib="NCI CGAP Co26"
 /note="Organ: colon; Vector: pAMP1; mRNA made from normal colonic mucosa, cDNA made by oligo-dT priming. Directionally cloned into UDG sites. Size-selected on agarose gel, average insert size 300 bp. Primary library."

FEATURES
 source

cDNA Library Preparation: David B. Krizman, Ph.D.
Reference: Krizman et al. (1996) Cancer Research
56:5380-5383."

ORIGIN
Query Match 47.8%; Score 11; DB 4; Length 50;
Best Local Similarity 100.0%; Pred. No. 3.2e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CATGACTGCAG 12
LOCUS
DEFINITION

RESULT 7

AW249907/c
LOCUS
DEFINITION

AW249907

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

NIH-MGC http://mgc.nci.nih.gov/.

National Institutes of Health, Mammalian Gene Collection (MGC)

Unpublished (1999)

Other ESTs: 2821659.5prime

Contact: Robert Strausberg, Ph.D.

Email: cgabs-r@mail.nih.gov

Tissue Procurement: DCTD/DTF cDNA Library Preparation: Ling

Hong/Rubin Laboratory cDNA Library Arrayed by: The I.M.A.G.E.

Consortium (LNL) DNA Sequencing by: Berkeley MGC sequencing

Project Clone distribution: MGC clone distribution information can

be found through the I.M.A.G.E. Consortium/LNL at:

www.bio.lnl.gov/bbrp/image/image.html Base Calling / Quality

Scores: PHRED from University of Washington Genome Center. Vector

Trimming: cross match from University of Washington Genome Center

PHRAP suite. Poly-T Identification: patMatch.pl from Berkeley

Drosophila Genome Project. University of Washington Genome Center:

http://www.genome.washington.edu Low Quality Sequence: 30

contiguous PHRED high quality bases following vector sequence. Very

Low Quality Sequence: Trace file contained 52 contiguous distinct

peaks following vector sequence. Polyadenylation: Based upon the

presence of a XhoI site followed by a run of 14 or more T residues

at the beginning of the sequence, this cDNA insert was

Polyadenylated.

Plate: L16M7 row: G column: 4

High quality sequence stop: 30.

Location/Qualifiers

1. 52

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:2821659"

/tissue_type="small cell carcinoma"

/cell_line="MGC3"

/lab_host="DH10B (phage-resistant)"

/clone_lib="NIH_MGC_7"

/notes="Organ: lung; Vector: pOTB7; Site 1: XhoI; Site 2:

EcoRI; cDNA made by oligo-dT priming. Directionally

cloned into EcoRI/XhoI sites using the following 5'

adaptor: GGACGAG(G). Size-selected >500bp for average

insert size 1.8kb. Library constructed by Ling Hong in

the laboratory of Gerald M. Rubin (University of

California, Berkeley) using ZAP-cDNA synthesis kit

(Stratagene) and Superscript II RT (Life Technologies)."

ORIGIN

Query Match

47.8%; Score 11; DB 2; Length 52;

Best Local Similarity 100.0%; Pred. No. 3.2e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ACTGCAATTC 16

|||||

Db 41 ACTGCAATTC 31

RESULT 8

CNS564600/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

CNS564600

tag20b04.x1 Hydra EST -Kiel 1 Hydra magnipapillata cDNA 3', mRNA

sequence.

CNS564600

CNS564600.1 GI:46973904

EST.

Hydra magnipapillata

Hydra magnipapillata

Eukaryota; Metazoa; Cnidaria; Hydrozoa; Anthomedusae;

Hydridae; Hydra;

1 (bases 1 to 55)

Bode,H., Blumberg,B., Steele,R., Wigge,P., Gee,L., Nguyen,Q.,

Martinez,D., Kibler,D., Hampson,S., Clifton,S., Pape,D., Marra,M.,

Hillier,L., Martin,J., Wylie,T., Dante,M., Theising,B., Bowers,Y.,

Gibbons,M., Ritter,E., Bennett,J., Ronko,I., Teagareishvili,R.,

Maguire,L., Kennedy,S., Waterston,R. and Wilson,R.

WashU Hydra EST Project

Unpublished (2002)

Contact: Hans Bode

WashU Hydra EST Project

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA

Tel: 314 286 1800

Fax: 314 286 1810

Email: est@watson.wustl.edu

Library was constructed by Konstantin Khalturin, Zoologisches

Institut, Univ. Kiel, Germany Library materials provided by Thomas

Borch, Zoologisches Institut, CAU Kiel, Germany DNA sequencing by:

Washington University Genome Sequencing Center For information on

obtaining a clone please contact: Hans Bode (hxbode@uci.edu)

Seq primer: degenerate primer.

Location/Qualifiers

1. 55

/organism="Hydra magnipapillata"

/mol_type="mRNA"

/db_xref="taxon:6085"

/lab_host="DH5a"

/clone_lib="Hydra EST -Kiel 1"

/note="Vector: pSPORT1; Site 1: Not I; Site 2: Sal I;

pSPORT 1 Vector is ampicillin resistant, M13 reverse

primer was used by us for sequencing of 5' parts of

inserts; 3' parts of cDNAs contain long polyA tracks which

makes sequencing from 3' direction complicated"

ORIGIN

Query Match

47.8%; Score 11; DB 7; Length 55;

Best Local Similarity 100.0%; Pred. No. 3.2e+04;

Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 13 TTCCGGTCTTT 23

|||||

Db 37 TTCCGGTCTTT 27

RESULT 9

BH918919/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

BH918919

BH918919.1 GI:22808353

GSS.

BH918919
3525.1_63.1_A10.2EL_x.1 3526 - RescueMu Grid K Zea mays genomic,
genomic survey sequence.
ACCESSION
VERSION
KEYWORDS
GSS.

```

SOURCE
ORGANISM
Zea mays
Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
1 (bases 1 to 57)
REFERENCE
AUTHORS
Walbot, V.
TITLE
Maize genomic sequences found using engineered RescueMu transposon
JOURNAL
Unpublished (2001)
COMMENT
Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Possible ligation site of ends cut by 2 different endonucleases.
Reverse complemented post-ligation sequence from source sequence.
Plate: 3526_1_63_1 row: 9
Class: transposon-tagged.
FEATURES
Location/Qualifiers
1..57
/organism="Zea mays"
/mol_type="genomic DNA"
/cultivar="mixed background W23/A188/B73"
/db_xref="taxon:4577"
/tissue_type="leaf"
/dev_stage="adult"
/lab_host="DH10B"
/clone_lib="3526 - RescueMu Grid K"
/notes="Organ: leaf; Vector: RescueMu (engineered from
pBluescript backbone); Site 1: BamHI; Site 2: BglII;
RescueMu is a 4.9 kb, modified maize Mu transposon
designed to allow plasmid rescue from total genomic DNA.
Mu elements insert preferentially into transcription
units. For more information on RescueMu, go to the web
site 'www.zmldb.iastate.edu' and follow the links for
'RescueMu.' Grid K was grown at Molokai, Hawaii in Winter
2000-2001. DNA was extracted from leaf punches, double
digested using BamHI and BglII, and ligated to form
circular plasmids. DH10B cells were transformed and then
screened on LB plates with ampicillin."
ORIGIN
Query Match 47.8%; Score 11; DB 8; Length 57;
Best Local Similarity 100.0%; Pred. No. 3.2e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CATGACTGCAAA 12
|||||
Db 51 CATGACTGCAAA 41

RESULT 10
AJ588850
LOCUS
Arabidopsis thaliana T-DNA flanking sequence, left border, clone
DEFINITION
358F07, genomic survey sequence.
ACCESSION
AJ588850
VERSION
AJ588850.1 GI:37938474
KEYWORDS
GSS; left border; T-DNA flanking sequence.
SOURCE
Arabidopsis thaliana (thale cress)
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosoids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1
REFERENCE
AUTHORS
Brunaud, V., Balzerque, S., Dubreucq, B., Aubourg, S., Samson, F.,
Chauvin, S., Bechtold, N., Cruaud, C., DeRose, R., Pelletier, G.,
Lepiniec, L., Caboche, M. and Lecharny, A.
TITLE
T-DNA integration into the Arabidopsis genome depends on sequences
of pre-insertion sites
JOURNAL
EMBO Rep. 3 (12), 1152-1157 (2002)

Zea mays
22363535
PUBMED
12446565
REFERENCE
2 (bases 1 to 19)
AUTHORS
Balzerque, S.
TITLE
Direct Submission
JOURNAL
Submitted (23-Oct-2003) Balzerque S., UMRGV, INRA/CNRS, 2 rue
Gaston Cremieux, 91057 Evry cedex, FRANCE
COMMENT
PCR was performed on DNA from transformants of Arabidopsis thaliana
plants from INRA (Versailles). The DNA fragment(s) resulting from
the PCR were directly sequenced from the left or the right border
to determine the genomic sequence flanking the insertion. T-DNA
derived sequences were removed. Information to order the
corresponding mutant line and a link to a database providing a
graphical display of the insertion site are available at
http://dbgap.versailles.inra.fr/publiclines/. This sequence has
been generated in the framework of the French plant genomics
program 'Genoplante' (http://www.genoplante.com and
http://genoplante-info.inbio.gen.fr).
FEATURES
Location/Qualifiers
1..19
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/cultivar="Wassillewskija"
/db_xref="taxon:3702"
/clone_lib="358F07"
/clone_lib="Arabidopsis thaliana T-DNA insertion lines"
/misc_feature
1..19
/notes="T-DNA flanking sequence
left border"
ORIGIN
Query Match 43.5%; Score 10; DB 9; Length 19;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 TCAGCTGCAAT 13
|||||
Db 3 TCAGCTGCAAT 12

RESULT 11
AG197078
LOCUS
Pan troglodytes DNA, clone: RP43-077A10.TJ, genomic survey
DEFINITION
sequence.
ACCESSION
AG197078
VERSION
AG197078.1 GI:45229254
KEYWORDS
GSS.
SOURCE
Pan troglodytes
ORGANISM
Pan troglodytes (chimpanzee)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
1
REFERENCE
AUTHORS
Park, H., Kim, Y., Kim, S., Han, Y., Woo, T., Park, K., Eun, C.J.,
Hoon, S.T., Chu, M., Kim, H., Joo, S., Kim, C., Song, W. and Yoo, H.
TITLE
BAC end sequences of Library RP-43
JOURNAL
Unpublished
REFERENCE
2 (bases 1 to 25)
AUTHORS
Park, H., Kim, Y., Kim, S., Han, Y., Woo, T., Park, K., Eun, C.J.,
Hoon, S.T., Chu, M., Kim, H., Joo, S., Kim, C., Song, W. and Yoo, H.
TITLE
Direct Submission
JOURNAL
Submitted (07-JAN-2002) Hong-Seog Park, Korea Research Institute of
Biotechnology and Biotechnology (KRIIB), Genome Research Center (GRC);
52, Oun-dong, Yuseong-gu, Daejeon 305-333, Korea
(E-mail: redstone@mail.kribb.re.kr, URL: http://phs.grc.kribb.re.kr/,
Tel: 82-42-866-7181, Fax: 82-42-860-4409)
COMMENT
Clones are derived from the chimpanzee BAC library RP-43 This BAC
end was generated during the R&D process and may have higher chance
of clone tracking errors.
PRIMERS
Sequencing: TJ
LIBRARY
Vector : pBACE3.6

```

```

R.Site 1 : EcORI
R.Site 2 : EcORI.
Location/Qualifiers
1. .25
/organism="Pan troglodytes"
/mol_type="genomic DNA"
/db_xref="taxon:9598"
/clone="RP43-077A10.TJ"
/sex="male"
/cell_type="lymphocytes"
/clone_lib="RP-43 Chimpanzee Male BAC Library"

ORIGIN
Query Match 43.5%; Score 10; DB 9; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 CTGCAATTCC 16
|||||
Db 1 CTGCAATTCC 10

RESULT 12
BX660142 34 bp DNA linear GSS 04-APR-2004
LOCUS Arabidopsis thaliana T-DNA flanking sequence GK-650H01-021296,
DEFINITION genomic survey sequence.
ACCESSION BX660142
VERSION BX660142.1 GI:37616530
KEYWORDS GSS.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1
Li, Y., Rosso, M.G., Strizhov, N., Viehoveer, P. and Weishaar, B.
GABI-Kat SimpleSearch: a flanking sequence tag (FST) database for
the identification of T-DNA insertion mutants in Arabidopsis
thaliana
JOURNAL Bioinformatics 19 (11), 1441-1442 (2003)
MEDLINE 22755829
PUBMED 12874060
REFERENCE 2
Rosso, M.G., Li, Y., Strizhov, N., Reiss, B., Dekker, K. and
Weishaar, B.
An Arabidopsis thaliana T-DNA mutagenized population (GABI-Kat) for
flanking sequence tag-based reverse genetics
Plant Mol. Biol. 53 (1-2), 247-259 (2003)
23117147
14756321
REFERENCE 3
Strizhov, N., Li, Y., Rosso, M.G., Viehoveer, P., Dekker, K.A. and
Weishaar, B.
High-throughput generation of sequence indexes from T-DNA
mutagenized Arabidopsis thaliana lines
Biotechniques 35 (6), 1164-1168 (2003)
14682050
4 (bases 1 to 34)
Li, Y., Strizhov, N., Rosso, M.G. and Weishaar, B.
Direct Submission
Submitted (31-MAR-2004) Weishaar B., Max-Planck-Institut fuer
Zuechtungsforchung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany
It indicates an insertion within the locus defined by BAC clone
T19E23. Details on the protocols used for generation of the
sequence are described in References 1-3. The sequences are
generated at the MPI for Plant Breeding Research in the context of
the GABI-Kat project. GABI-Kat is part of the German Plant Genomics
program designated 'GABI'. Information on line availability can be
found at: http://www.mpiz-koeln.mpg.de/GABI-Kat/.
Location/Qualifiers
1. .34

FEATURES
source
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/db_xref="taxon:3702"
/clone="GK-650H01-021296"
/clone_lib="Arabidopsis thaliana T-DNA insertion lines"
/ecotype="Col-0"
/note="PCR was performed on DNA from Arabidopsis thaliana
plants (T1) which were transformed with the T-DNA from
vector PAC161 (GenBank accession number: AJ537514). The
lines contain one or more T-DNA insertions. The DNA
fragment(s) resulting from the PCR were directly sequenced
to determine the genomic sequence flanking the insertion.
T-DNA derived sequences were removed."

ORIGIN
Query Match 43.5%; Score 10; DB 9; Length 34;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 TGACTGCAAT 13
|||||
Db 17 TGACTGCAAT 26

RESULT 13
BX810737 36 bp DNA linear GSS 02-MAY-2002
LOCUS SALK 051126 Arabidopsis thaliana TDNA insertion lines Arabidopsis
DEFINITION thaliana genomic clone SALK_051126, genomic survey sequence.
ACCESSION BX810737
VERSION BX810737.1 GI:20388555
KEYWORDS GSS.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 36)
Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R.,
Gadrinab, C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prednis, L.,
Shinn, P., Zimmerman, J. and Ecker, J.R.
A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
Unpublished (2001)
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGNAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of
TDNA.
Class: TDNA tagged.
Location/Qualifiers
1. .36
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/ecotype="Col-0"
/db_xref="taxon:3702"
/clone="SALK_051126"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"

ORIGIN
Query Match 43.5%; Score 10; DB 8; Length 36;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;

```

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 14 TCCGGTCTTT 23
|||||
Db 18 TCCGGTCTTT 9

RESULT 14

DMES46528/c

LOCUS

DEFINITION

PI(RS5)5-HA-1904, clone library P[RS], genomic survey sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

source

misc_feature

ORIGIN

Query Match

Best Local Similarity

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 6 ACTGCAATTC 15
|||||
Db 23 ACTGCAATTC 14

RESULT 15

AA972482

LOCUS

DEFINITION

IMAGE:1579493 3' similar to TR:Q13526 Q13526 PIN1. ;, mRNA

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 20:30:00 ; Search time 233 Seconds
(without alignments)
584.352 Million cell updates/sec

Title: US-10-729-421-8
Perfect score: 23
Sequence: 1 tcatgactgaattccggtcttt 23

Scoring table: OLIGO NUC
Gapop 60.0 , Gapext 60.0

Searched: 4390206 seqs, 2959870667 residues

Word size : 10

Total number of hits satisfying chosen parameters: 684

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Listing first 45 summaries

Database : N_Geneseq_16Dec04:*
1: Geneseqn1980s:*
2: Geneseqn1990s:*
3: Geneseqn2000s:*
4: Geneseqn2001as:*
5: Geneseqn2001bs:*
6: Geneseqn2002as:*
7: Geneseqn2002bs:*
8: Geneseqn2003as:*
9: Geneseqn2003bs:*
10: Geneseqn2003cs:*
11: Geneseqn2003ds:*
12: Geneseqn2004as:*
13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	23	100.0	23	12	ADQ30638
2	23	100.0	46	12	ADQ30655
3	20	87.0	36	8	ABZ25451
4	20	87.0	36	9	AAL55873
5	20	87.0	36	12	ADM16871
6	20	87.0	39	8	ABZ25450
7	20	87.0	39	9	AAL55872
8	20	87.0	39	12	ADM16870
9	20	87.0	51	10	ADC06634
10	20	87.0	51	10	ADC06633
11	20	87.0	58	12	ADM16878
12	19	82.6	33	8	ABZ25440
13	19	82.6	33	9	AAL55862
14	19	82.6	33	12	ADM16860
15	19	82.6	36	8	ABZ25431
16	19	82.6	36	9	AAL57662
17	19	82.6	36	12	ADM16851
18	19	82.6	51	10	ADC06635
19	19	82.6	56	4	AAD14623
20	17	73.9	17	6	ACN09412

c	21	17	73.9	17	6	ACN00067
c	22	17	73.9	17	6	ACN13570
c	23	17	73.9	17	6	ACN03509
c	24	17	73.9	17	6	ACN14150
c	25	17	73.9	17	6	ACN05528
c	26	17	73.9	17	6	ACN12169
c	27	17	73.9	17	6	ACN04748
c	28	17	73.9	17	6	ACN09413
c	29	16	69.6	17	6	ACN12168
c	30	16	69.6	17	6	ACN05527
c	31	16	69.6	17	6	ACN12170
c	32	16	69.6	17	6	ACN01482
c	33	16	69.6	30	10	ADC06605
c	34	15	65.2	17	6	ACN03510
c	35	15	65.2	17	6	ACN15245
c	36	15	65.2	17	6	ACN01481
c	37	14	60.9	17	6	ACN14149
c	38	14	60.9	17	6	ACN13571
c	39	13	56.5	17	6	ACN00068
c	40	13	56.5	17	6	ACN04747
c	41	13	56.5	17	6	ACN15244
c	42	13	56.5	25	9	ACK26449
c	43	13	56.5	30	10	ADC06609
c	44	13	56.5	30	10	ADC06611
c	45	12	52.2	17	6	ACN09414

ALIGNMENTS

RESULT 1
ADQ30638
ID ADQ30638 standard; DNA; 23 BP.
XX
AC ADQ30638;
XX
DT 23-SEP-2004 (first entry)
XX
DE West Nile Virus capture oligonucleotide WNVVC8.
XX
KW ss; capture oligonucleotide; West Nile Virus; diagnosis.
XX
OS West Nile virus.
XX
PN WO2004055159-A2.
XX
PD 01-JUL-2004.
XX
PF 05-DEC-2003; 2003WO-US038750.
XX
PR 12-DEC-2002; 2002US-0432850P.
PR 20-JUN-2003; 2003US-0480431P.
XX (CHIR) CHIRON CORP.

PA Shyamala V;
XX
DR WPI; 2004-488058/46.
XX

PT New isolated oligonucleotides for accurately diagnosing West Nile virus infection or for capturing, detecting and quantitating West Nile virus in blood samples.
PT
XX

PS Claim 1; SEQ ID NO 8; 56pp; English.
XX

CC The invention relates to an isolated oligonucleotide not more than 60 nucleotides in length comprising a nucleotide sequence (S1) of at least 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g. CC 20, 21 or 23 bp) given in the specification derived from the West Nile virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence identity to the nucleotide sequence of (S1), or complements of (S1) and (S2). The oligonucleotide further comprises a detectable label at the 5'-end and/or the 3'-end. The detectable label is a fluorescent label
CC

CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
CC composition and methods are useful for accurately diagnosing West Nile
CC virus infection or for capturing, detecting and quantitating West Nile
CC virus in biological samples, particularly blood samples. This sequence
CC corresponds to a capture oligonucleotide of the invention.

XX Sequence 23 BP; 4 A; 6 C; 4 G; 9 T; 0 U; 0 Other;

Query Match 100.0%; Score 23; DB 12; Length 23;
Best Local Similarity 100.0%; Pred. No. 0.00091;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCATGACTGCAATTCGGTCTTT 23
DB 1 TCATGACTGCAATTCGGTCTTT 23

RESULT 2

ADQ30655
ID ADQ30655 standard; DNA; 46 BP.

XX AC ADQ30655;

DT 23-SEP-2004 (first entry)

XX West Nile Virus capture oligonucleotide poly-A-WNVVC8.

DE ss; capture oligonucleotide; West Nile Virus; diagnosis.

KW West Nile virus.

OS WO2004055159-A2.

XX 01-JUL-2004.

XX 05-DEC-2003; 2003WO-US038750.

XX 12-DEC-2002; 2002US-0432850P.

PR 20-JUN-2003; 2003US-0480431P.

XX (CHIR) CHIRON CORP.

PA Shyamala V;

XX WPI; 2004-488058/46.

XX New isolated oligonucleotides for accurately diagnosing West Nile virus
PT infection or for capturing, detecting and quantitating West Nile virus in
PT blood samples.

PS Example 1; SEQ ID NO 25; 56pp; English.

XX The invention relates to an isolated oligonucleotide not more than 60
CC nucleotides in length comprising a nucleotide sequence (S1) of at least
CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
CC 20, 21 or 23 bp) given in the specification derived from the West Nile
CC virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
CC identity to the nucleotide sequence of (S1), or complements of (S1) and
CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
CC end and/or the 3'-end. The detectable label is a fluorescent label
CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
CC composition and methods are useful for accurately diagnosing West Nile
CC virus infection or for capturing, detecting and quantitating West Nile
CC virus in biological samples, particularly blood samples. This sequence
CC corresponds to a capture oligonucleotide of the invention.

XX Sequence 46 BP; 27 A; 6 C; 4 G; 9 T; 0 U; 0 Other;

Query Match 100.0%; Score 23; DB 12; Length 46;
Best Local Similarity 100.0%; Pred. No. 0.00088;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCATGACTGCAATTCGGTCTTT 23
DB 24 TCATGACTGCAATTCGGTCTTT 46

RESULT 3

ABZ25451/c
ID ABZ25451 standard; DNA; 36 BP.

XX AC ABZ25451;

DT 27-MAR-2003 (first entry)

XX PCR primer FC117, SEQ ID 29.

XX Virucide; vaccine; horse; dog; cat; cattle; pig; bird; West Nile virus;
KW WNV; PCR; primer; ss.

XX Synthetic.

XX WO200281621-A2.

XX 17-OCT-2002.

XX 05-APR-2002; 2002WO-FR001200.

XX 06-APR-2001; 2001FR-00004737.

XX (MERI-) MERIAL.

XX Loosmore SM, Audonnet JF;

XX WPI; 2003-111799/10.

XX Vaccine for treatment or prevention of West Nile virus (WNV) infection,
PT for use in veterinary medicine, comprises a recombinant virus expressing
PT a WNV structural protein.

XX Example 18; Page 41; 56pp; French.

XX The present invention relates to a vaccine for protecting horses, dogs,
CC cats, cattle, pigs and birds against West Nile virus (WNV). The vaccine
CC comprises: (i) one or more recombinant avipox, NYVAC or WVA viruses that
CC express one of the WNV proteins prM, M and E and (ii) a vehicle or
CC excipient. The present sequence is a PCR primer, which was used in an
CC example from the invention

XX Sequence 36 BP; 8 A; 7 C; 10 G; 11 T; 0 U; 0 Other;

Query Match 87.0%; Score 20; DB 8; Length 36;
Best Local Similarity 100.0%; Pred. No. 0.052;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCATGACTGCAATTCGGTCTTC 20
DB 31 TCATGACTGCAATTCGGTCTTC 12

RESULT 4

AAL55873/c
ID AAL55873 standard; DNA; 36 BP.

XX AC AAL55873;

DT 06-NOV-2003 (first entry)

XX FC117 PCR primer used to amplify the plasmid pFC115.

XX Immunogenic composition; West Nile fever virus; WNV; prM; M; membrane; E;
KW pre-membrane protein; envelope; virucide; vaccine; FC117; primer; PCR;
KW ss; plasmid pFC115.

OS Unidentified.
 XX US2003104008-A1.
 XX
 XX 05-JUN-2003.
 XX
 XX 04-APR-2002; 2002US-00116298.
 XX
 XX 06-APR-2001; 2001US-0281923P.
 XX
 XX (LOOS/) LOOSMORE S M.
 XX (AUDO/) AUDONNET J F.
 XX
 XX Loosmore SM, Audonnet JF;
 XX WPI; 2003-567944/53.
 XX
 XX New immunogenic composition comprising a recombinant avipox virus that
 PT expresses in vivo in the animal the West Nile (WN) proteins prM, M or E,
 PT useful for inducing an immunological response against WN virus.
 XX
 XX Example 18; Page 14; 24pp; English.
 XX
 XX The invention relates to a novel immunogenic composition for inducing an
 CC immune response against West Nile fever virus (WNV) in an animal. The
 CC composition comprises a vehicle or excipient and a recombinant avipox
 CC virus that expresses in vivo in the animal the WNV proteins prM (pre-
 CC membrane protein), M (membrane protein) or E (envelope protein). The
 CC animal is selected from canine, feline, bovine, porcine, chicken, equine,
 CC a duck, a goose or a turkey. The composition of the invention
 CC demonstrates virucide activity and may be useful as a vaccine against
 CC WNV. The current sequence is that of the FC117 PCR primer of the
 CC invention which was used to amplify the plasmid pFC115
 XX
 XX Sequence 36 BP; 8 A; 7 C; 10 G; 11 T; 0 U; 0 Other;
 SQ
 Query Match 87.0%; Score 20; DB 9; Length 36;
 Best Local Similarity 100.0%; Pred. No. 0.052;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TCATGACTGCAATTCGGTC 20
 DB 31 TCATGACTGCAATTCGGTC 12
 RESULT 5
 ADM16871/c
 ID ADM16871 standard; DNA; 36 BP.
 XX
 XX ADM16871;
 XX
 XX 20-MAY-2004 (first entry)
 XX
 XX Plasmid pFC115 PCR primer #1.
 DE
 XX Immunogen; vaccine; West Nile virus; ss; PCR; primer.
 XX
 XX Synthetic.
 OS
 XX US2004037848-A1.
 XX
 XX 26-FEB-2004.
 XX
 XX 26-FEB-2003; 2003US-00374953.
 XX
 XX 06-APR-2001; 2001US-0281923P.
 XX 04-APR-2002; 2002US-00116298.
 XX
 XX (AUDO/) AUDONNET J F.
 XX (MINK/) MINKE J M.
 XX (LOOS/) LOOSMORE S M.
 XX (KARA/) KARACA K.
 XX

PI Audonnet JF, Minke JM, Loosmore SM, Karaca K;
 XX WPI; 2004-191012/18.
 XX
 XX Vaccine composition, useful in inducing an immune response against West
 PT Nile virus, comprises a vector that contains heterologous nucleic acid
 PT molecule(s), and that expresses in vivo in the animal a WNV protein.
 XX
 XX Example 18; SEQ ID NO 29; 36pp; English.
 XX
 XX The invention relates to an immunogenic or vaccine composition which
 CC induces an immune response against West Nile virus (WNV) in an animal
 CC susceptible to WNV comprises a vector that contains heterologous nucleic
 CC acid molecule(s) and that expresses in vivo in the animal a WNV E; WNV
 CC prM and E; WNV M and E; WNV prM, WNV M and E, WNV polypeptide prM-E, WNV
 CC polypeptide M-E, or WNV polypeptide prM-W-E. The composition is useful
 CC for inducing an immunological or protective immune response against WNV
 CC and against another pathogen of the animal. Also inducing an
 CC immunological or protective immune response against WNV in an animal
 CC comprises administering to the animal (a) the immunogenic or vaccine
 CC composition and (b) a WNV isolated antigen, immunogen or epitope, where
 CC (a) is administered prior to (b) in a prime-boost regimen, or (b) is
 CC administered prior to (a) in a prime-boost regimen, or (a) and (b) are
 CC administered together, either sequentially or in admixture. The present
 CC sequence is used in the exemplification of the invention.
 XX
 XX Sequence 36 BP; 8 A; 7 C; 10 G; 11 T; 0 U; 0 Other;
 SQ
 Query Match 87.0%; Score 20; DB 12; Length 36;
 Best Local Similarity 100.0%; Pred. No. 0.052;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TCATGACTGCAATTCGGTC 20
 DB 31 TCATGACTGCAATTCGGTC 12
 RESULT 6
 ABZ25450/c
 ID ABZ25450 standard; DNA; 39 BP.
 XX
 XX AC ABZ25450;
 XX
 XX 27-MAR-2003 (first entry)
 DT
 XX
 XX West Nile Virus PCR primer FC116, SEQ ID 28.
 DE
 XX Virucide; vaccine; horse; dog; cat; cattle; pig; bird; West Nile virus;
 KW WNV; PCR; primer; ss.
 XX
 XX West Nile Virus.
 OS
 XX WO200281621-A2.
 PN
 XX 17-OCT-2002.
 PD
 XX 05-APR-2002; 2002WO-PR001200.
 PF
 XX 06-APR-2001; 2001FR-00004737.
 PR
 XX (MERI-) MERIAL.
 XX
 XX Loosmore SM, Audonnet JF;
 PI WPI; 2003-111799/10.
 XX
 XX Vaccine for treatment or prevention of West Nile virus (WNV) infection,
 PT for use in veterinary medicine, comprises a recombinant virus expressing
 PT a WNV structural protein.
 XX
 XX Example 17; Page 40; 56pp; French.
 PS
 XX The present invention relates to a vaccine for protecting horses, dogs,
 CC

CC cats, cattle, pigs and birds against West Nile virus (WNV). The vaccine
 CC comprises: (i) one or more recombinant avipox, NVAC or MVA viruses that
 CC express one of the WNV proteins prM, M and E and (ii) a vehicle or
 CC excipient. The present sequence is a PCR primer, which was used in an
 CC example from the invention

XX
 SQ Sequence 39 BP; 9 A; 6 C; 9 G; 15 T; 0 U; 0 Other;

Query Match 87.0%; Score 20; DB 8; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.052;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TCATGACTGCAATTCGGGTC 20
 DB 34 TCATGACTGCAATTCGGGTC 15

RESULT 7
 AAL55872/c
 ID AAL55872 standard; DNA; 39 BP.
 XX
 AC AAL55872;
 XX
 DT 06-NOV-2003 (first entry)
 XX
 DE FC116 RT-PCR primer used to amplify West Nile fever virus RNA.
 XX
 KW Immunogenic composition; West Nile fever virus; WNV; prM; M; membrane; E;
 KW pre-membrane protein; envelope; virucide; vaccine; FC116; RT-PCR; primer;
 KW PCR; ss.
 XX
 OS Unidentified.
 OS Synthetic.
 XX
 PN US2003104008-A1.
 XX
 PD 05-JUN-2003.
 XX
 PF 04-APR-2002; 2002US-00116298.
 XX
 PR 06-APR-2001; 2001US-0281923P.
 XX
 PA (LOOS/) LOOSMORE S M.
 PA (AUDO/) AUDONNET J F.
 XX
 PI Loosmore SM, Audonnet JF;
 XX
 XX WPI; 2003-567944/53.
 XX
 PT New immunogenic composition comprising a recombinant avipox virus that
 PT expresses in vivo in the animal the West Nile (WN) proteins prM, M or E,
 PT useful for inducing an immunological response against WN virus.
 XX
 PS Example 17; Page 14; 24pp; English.
 XX
 CC The invention relates to a novel immunogenic composition for inducing an
 CC immune response against West Nile fever virus (WNV) in an animal. The
 CC composition comprises a vehicle or excipient and a recombinant avipox
 CC virus that expresses in vivo in the animal the WNV proteins prM (pre-
 CC membrane protein), M (membrane protein) or E (envelope protein). The
 CC animal is selected from canine, feline, bovine, porcine, chicken, equine,
 CC a duck, a goose or a turkey. The composition of the invention
 CC demonstrates virucide activity and may be useful as a vaccine against
 CC WNV. The current sequence is that of the FC116 RT-PCR primer of the
 CC invention which was used to amplify West Nile fever virus RNA

XX
 SQ Sequence 39 BP; 9 A; 6 C; 9 G; 15 T; 0 U; 0 Other;

Query Match 87.0%; Score 20; DB 9; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.052;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TCATGACTGCAATTCGGGTC 20

DB 34 TCATGACTGCAATTCGGGTC 15

RESULT 8
 ADM16870/c
 ID ADM16870 standard; DNA; 39 BP.
 XX
 AC ADM16870;
 XX
 DT 20-MAY-2004 (first entry)
 XX
 DE West Nile virus RT-PCR primer #10.
 XX
 KW immunogen; vaccine; West Nile virus; ss; reverse transcriptase; RT-PCR;
 KW primer.
 XX
 OS West Nile virus.
 XX
 PN US2004037848-A1.
 XX
 PD 26-FEB-2004.
 XX
 PF 26-FEB-2003; 2003US-00374953.
 XX
 PR 06-APR-2001; 2001US-0281923P.
 PR 04-APR-2002; 2002US-00116298.
 XX
 PA (AUDO/) AUDONNET J F.
 PA (MINK/) MINK J M.
 PA (LOOS/) LOOSMORE S M.
 PA (KARA/) KARACA K.
 XX
 PI Audonnet JF, Minke JM, Loosmore SM, Karaca K;
 XX
 XX WPI; 2004-191012/18.
 XX
 PT Vaccine composition, useful in inducing an immune response against West
 PT Nile virus, comprises a vector that contains heterologous nucleic acid
 PT molecule(s), and that expresses in vivo in the animal a WNV protein.
 XX
 PS Example 17; SEQ ID NO 28; 36pp; English.
 XX
 CC The invention relates to an immunogenic or vaccine composition which
 CC induces an immune response against West Nile virus (WNV) in an animal
 CC susceptible to WNV comprises a vector that contains heterologous nucleic
 CC acid molecule(s) and that expresses in vivo in the animal a WNV E; WNV
 CC prM and E; WNV M and E; WNV prM, WNV M and E, WNV polypeptide prM-E, WNV
 CC polypeptide M-E, or WNV polypeptide prM-M-E. The composition is useful
 CC for inducing an immunological or protective immune response against WNV
 CC and against another pathogen of the animal. Also inducing an
 CC immunological or protective immune response against WNV in an animal
 CC comprises administering to the animal (a) the immunogenic or vaccine
 CC composition and (b) a WNV isolated antigen, immunogen or epitope, where
 CC (a) is administered prior to (b) in a prime-boost regimen, or (b) is
 CC administered prior to (a) in a prime-boost regimen, or (a) and (b) are
 CC administered together, either sequentially or in admixture. The present
 CC sequence is used in the exemplification of the invention.

XX
 SQ Sequence 39 BP; 9 A; 6 C; 9 G; 15 T; 0 U; 0 Other;

Query Match 87.0%; Score 20; DB 12; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.052;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TCATGACTGCAATTCGGGTC 20
 DB 34 TCATGACTGCAATTCGGGTC 15

RESULT 9
 ADC06634/c
 ID ADC06634 standard; DNA; 51 BP.

```

XX AC ADC06634;
XX AC
XX DT 18-DEC-2003 (first entry)
XX DE PCR primer SEQ ID 35 used during construction of WNV/DEN4 chimeras.
XX KW West Nile virus; WNV; DEN4; Dengue virus type 4; virucide; vaccine; ss;
XX KW PCR; primer.
XX OS Unidentified.
XX PN WO2003059384-A1.
XX PD 24-JUL-2003.
XX XX
XX XX 09-JAN-2003; 2003WO-US000594.
XX PF 10-JAN-2002; 2002US-0347281P.
XX PR (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX PA Pletnev AG, Putnak JR, Chanock RM, Murphy BR, Whitehead SS;
XX PI Blaney JE;
XX XX
XX WPI; 2003-636686/60.
XX XX
XX PT Novel nucleic acid chimera comprising nucleic acids encoding structural
XX PT protein from West Nile virus and non-structural proteins from wild-type
XX PT strain of dengue virus useful for producing live West Nile virus
XX PT vaccines.
XX XX
XX PS Disclosure; Page 20; 53pp; English.
XX CC The invention relates to a novel nucleic acid chimera comprising a first
XX CC nucleotide sequence encoding at least one structural protein from a West
XX CC Nile virus (WNV) and a second nucleotide sequence encoding non-structural
XX CC proteins from a wild-type strain of Dengue virus (DEN), such as Dengue
XX CC virus type 4 (DEN4). The nucleotide of the invention demonstrates
XX CC virucide activity and may be useful for producing a WNV vaccine. The
XX CC current sequence is that of the PCR primer of the invention which was
XX CC used during the construction of the WNV/DEN4 chimeras.
XX XX
XX SQ Sequence 51 BP; 20 A; 9 C; 13 G; 9 T; 0 U; 0 Other;
Query Match 87.0%; Score 20; DB 10; Length 51;
Best Local Similarity 100.0%; Pred. No. 0.051;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCATGACTGCAATTCCGGTC 20
DB 46 TCATGACTGCAATTCCGGTC 27
|||||
XXXXXXXXXXXXX
RESULT 10
ADC06633/c
ID ID ADC06633 standard; DNA; 51 BP.
XX AC ADC06633;
XX XX
XX DT 18-DEC-2003 (first entry)
XX DE PCR primer SEQ ID 34 used during construction of WNV/DEN4 chimeras.
XX XX
XX KW West Nile virus; WNV; DEN4; Dengue virus type 4; virucide; vaccine; ss;
XX KW PCR; primer.
XX OS Unidentified.
XX PN WO2003059384-A1.
XX PD 24-JUL-2003.
XX XX
XX XX

```

XX PS Example 28; SEQ ID NO 36; 36pp; English.

XX CC The invention relates to an immunogenic or vaccine composition which induces an immune response against West Nile virus (WNV) in an animal susceptible to WNV comprising a vector that contains heterologous nucleic acid molecule(s) and that expresses in vivo in the animal a WNV E; WNV prM and E; WNV M and E; WNV prM, WNV M and E, WNV polyprotein prM-E, WNV polyprotein M-E, or WNV polyprotein prM-M-E. The composition is useful for inducing an immunological or protective immune response against WNV and against another pathogen of the animal. Also inducing an immunological or protective immune response against WNV in an animal comprises administering to the animal (a) the immunogenic or vaccine composition and (b) a WNV isolated antigen, immunogen or epitope, where (a) is administered prior to (b) in a prime-boost regimen, or (b) is administered prior to (a) in a prime-boost regimen, or (a) and (b) are administered together, either sequentially or in admixture. The present sequence is used in the exemplification of the invention.

XX SQ Sequence 58 BP; 13 A; 10 C; 14 G; 21 T; 0 U; 0 Other;

Query Match 87.0%; Score 20; DB 12; Length 58;
Best Local Similarity 100.0%; Pred. No. 0.051;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCATGACTGCAATTCGGTC 20
DB 53 TCATGACTGCAATTCGGTC 34

RESULT 12
AB225440/c
ID AB225440 standard; DNA; 33 BP.

XX AC AB225440;

XX DT 27-MAR-2003 (first entry)

XX DE PCR primer FC110, SEQ ID 18.

XX KW Virucide; vaccine; horse; dog; cat; cattle; pig; bird; West Nile virus; WNV; PCR; primer; ss.

XX OS Synthetic.

XX PN WO20281621-A2.

XX PD 17-OCT-2002.

XX PF 05-APR-2002; 2002WO-FR001200.

XX PR 06-APR-2001; 2001FR-00004737.

XX PA (MERI-) MERIAL.

XX PI Loosmore SM, Audonnet JF;

XX PS WPI; 2003-111799/10.

XX PT Vaccine for treatment or prevention of West Nile virus (WNV) infection, for use in veterinary medicine, comprises a recombinant virus expressing a WNV structural protein.

XX PS Example 9; Page 34; 56pp; French.

XX CC The present invention relates to a vaccine for protecting horses, dogs, cats, cattle, pigs and birds against West Nile virus (WNV). The vaccine comprises: (i) one or more recombinant avipox, NYVAC or MVA viruses that express one of the WNV proteins prM, M and E and (ii) a vehicle or excipient. The present sequence is a PCR primer, which was used in an example from the invention

XX SQ Sequence 33 BP; 7 A; 7 C; 9 G; 10 T; 0 U; 0 Other;

Query Match 82.6%; Score 19; DB 8; Length 33;
Best Local Similarity 100.0%; Pred. No. 0.2;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCATGACTGCAATTCGGT 19
DB 28 TCATGACTGCAATTCGGT 10

RESULT 13
AAL55862/c
ID AAL55862 standard; DNA; 33 BP.

XX AC AAL55862;

XX DT 06-NOV-2003 (first entry)

XX DE FC110 PCR primer used to amplify the plasmid pFC105.

XX KW Immunogenic composition; West Nile fever virus; WNV; prM; M; membrane; E; pre-membrane protein; envelope; virucide; vaccine; FC110; primer; PCR; ss; plasmid pFC105.

XX OS Unidentified.

XX PN US2003104008-A1.

XX PD 05-JUN-2003.

XX PF 04-APR-2002; 2002US-00116298.

XX PR 06-APR-2001; 2001US-0281923P.

XX PA (LOOS/) LOOSMORE S M.

XX PI (AUDO/) AUDONNET J F.

XX PI Loosmore SM, Audonnet JF;

XX DR WPI; 2003-567944/53.

XX PT New immunogenic composition comprising a recombinant avipox virus that expresses in vivo in the animal the West Nile (WN) proteins prM, M or E, useful for inducing an immunological response against WN virus.

XX PS Example 10; Page 12; 24pp; English.

XX CC The invention relates to a novel immunogenic composition for inducing an immune response against West Nile fever virus (WNV) in an animal. The composition comprises a vehicle or excipient and a recombinant avipox virus that expresses in vivo in the animal the WNV proteins prM (pre-membrane protein), M (membrane protein) or E (envelope protein). The animal is selected from canine, feline, bovine, porcine, chicken, equine, a duck, a goose or a turkey. The composition of the invention demonstrates virucide activity and may be useful as a vaccine against WNV. The current sequence is that of the FC110 PCR primer of the invention which was used to amplify the plasmid pFC105

XX SQ Sequence 33 BP; 7 A; 7 C; 9 G; 10 T; 0 U; 0 Other;

Query Match 82.6%; Score 19; DB 9; Length 33;
Best Local Similarity 100.0%; Pred. No. 0.2;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCATGACTGCAATTCGGT 19
DB 28 TCATGACTGCAATTCGGT 10

RESULT 14
ADM16860/c
ID ADM16860 standard; DNA; 33 BP.

XX

```
AC ADM16860;
XX
XX 20-MAY-2004 (first entry)
XX
XX Plasmid pFC 105 PCR primer #1.
XX
XX immunogen; vaccine; West Nile virus; ss; PCR; primer.
XX
XX Synthetic.
XX
XX OS US2004037848-A1.
XX
XX PN 26-FEB-2004.
XX
XX PD
XX
XX PF 26-FEB-2003; 2003US-00374953.
XX
XX PR 06-APR-2001; 2001US-0281923P.
XX
XX BR 04-APR-2002; 2002US-00116298.
XX
XX
XX (AUO/) AUDONNET J F.
XX (MINK/) MINKE J M.
XX PA (LOOS/) LOOSMORE S M.
XX PA (KARA/) KARACA K.
XX
XX PI Audonnet JF, Minke JM, Loosmore SM, Karaca K;
XX
XX WIPI; 2004-191012/18.
XX
XX Vaccine composition, useful in inducing an immune response against West
XX Nile virus, comprises a vector that contains heterologous nucleic acid
XX molecule(s), and that expresses in vivo in the animal a WNV protein.
XX
XX Example 10; SEQ ID NO 18; 36pp; English.
XX
XX The invention relates to an immunogenic or vaccine composition which
XX induces an immune response against West Nile virus (WNV) in an animal
XX susceptible to WNV comprises a vector that contains heterologous nucleic
XX acid molecule(s) and that expresses in vivo in the animal a WNV E; WNV
XX prM and E; WNV M and E; WNV prM, WNV M and E, WNV polyprotein prM-E, WNV
XX polyprotein M-E, or WNV polyprotein prM-M-E. The composition is useful
XX for inducing an immunological or protective immune response against WNV
XX and against another pathogen of the animal. Also inducing an
XX immunological or protective immune response against WNV in an animal
XX comprises administering to the animal (a) the immunogenic or vaccine
XX composition and (b) a WNV isolated antigen, immunogen or epitope, where
XX (a) is administered prior to (b) in a prime-boost regimen, or (b) is
XX administered prior to (a) in a prime-boost regimen, or (a) and (b) are
XX administered together, either sequentially or in admixture. The present
XX sequence is used in the exemplification of the invention.
XX
XX Sequence 33 BP; 7 A; 7 C; 9 G; 10 T; 0 U; 0 Other;
XX
Query Match 82.6%; Score 19; DB 12; Length 33;
Best Local Similarity 100.0%; Pred. No. 0.2;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCATGACTGCAATTCCGGT 19
DB 28 TCATGACTGCAATTCCGGT 10
RESULT 15
ABZ25431/c
ID ABZ25431 standard; DNA; 36 BP.
XX
XX AC ABZ25431;
XX
XX DT 27-MAR-2003 (first entry)
XX
XX DE West Nile Virus PCR primer FC107, SEQ ID 9.
XX
XX Virucide; vaccine; horse; dog; cat; cattle; pig; bird; West Nile virus;
XX WNV; PCR; primer; ss.
```

```
XX
XX OS West Nile Virus.
XX
XX PN WO200281621-A2.
XX
XX PD 17-OCT-2002.
XX
XX PF 05-APR-2002; 2002WO-FR001200.
XX
XX PR 06-APR-2001; 2001FR-00004737.
XX
XX PA (MERI-) MERIAL.
XX
XX PI Loosmore SM, Audonnet JF;
XX
XX WIPI; 2003-111799/10.
XX
XX Vaccine for treatment or prevention of West Nile virus (WNV) infection,
XX for use in veterinary medicine, comprises a recombinant virus expressing
XX a WNV structural protein.
XX
XX Example 7; Page 31; 56pp; French.
XX
XX The present invention relates to a vaccine for protecting horses, dogs,
XX cats, cattle, pigs and birds against West Nile virus (WNV). The vaccine
XX comprises: (i) one or more recombinant avipox, NVVAC or MVA viruses that
XX express one of the WNV proteins prM, M and E and (ii) a vehicle or
XX excipient. The present sequence is a PCR primer, which was used in an
XX example from the invention
XX
XX Sequence 36 BP; 8 A; 6 C; 8 G; 14 T; 0 U; 0 Other;
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Best Local Similarity 100.0%; Pred. No. 0.2;
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DB 31 TCATGACTGCAATTCCGGT 13
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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C 1	20	87.0	36	6	AX589722 Sequence
C 2	20	87.0	39	6	AX589721 Sequence
C 3	19	82.6	33	6	AX589711 Sequence
C 4	19	82.6	36	6	AX589702 Sequence
C 5	19	82.6	56	6	AX224249 Sequence
C 6	12	52.2	25	6	AX182204 Sequence
7	12	52.2	25	6	AX382013 Sequence
8	12	52.2	34	6	BD173847 JNK inhib
9	12	52.2	51	6	AR444320 Sequence
10	12	52.2	51	6	AR444321 Sequence
11	12	52.2	60	6	CQ536043 Sequence
C 12	12	52.2	60	14	POLDIPI
C 13	11	47.8	15	6	AR119503 Sequence
C 14	11	47.8	16	6	AR285636 Sequence
C 15	11	47.8	16	6	AR397627 Sequence
16	11	47.8	20	6	AR167035 Sequence
17	11	47.8	20	6	AR210690 Sequence
C 18	11	47.8	20	6	AR301418 Sequence
C 19	11	47.8	20	6	AR313575 Sequence

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21	11	47.8	21	6	AR166997 Sequence
22	11	47.8	21	6	AR210652 Sequence
23	11	47.8	22	6	CQ768639 Sequence
24	11	47.8	24	6	AX935032 Sequence
C 25	11	47.8	25	6	AR364418 Sequence
C 26	11	47.8	25	6	AR568240 Sequence
C 27	11	47.8	26	6	AR542624 Sequence
C 28	11	47.8	26	6	AX235895 Sequence
29	11	47.8	26	6	AX402749 Sequence
C 30	11	47.8	27	6	BD183051 Nucleic a
C 31	11	47.8	27	6	I22149 Sequence 8
C 32	11	47.8	28	6	I13959 Sequence 38
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35	11	47.8	33	6	AR004393 Sequence
C 36	11	47.8	33	6	AR005205 Sequence
37	11	47.8	33	6	AR005206 Sequence
38	11	47.8	33	6	AR064955 Sequence
C 39	11	47.8	33	6	AR072936 Sequence
C 40	11	47.8	33	6	AR072938 Sequence
41	11	47.8	33	6	AR097185 Sequence
42	11	47.8	33	6	AR130683 Sequence
43	11	47.8	33	6	AR172032 Sequence
44	11	47.8	33	6	BD189149 HCV Genom
45	11	47.8	33	6	BD189296 HCV Genom

ALIGNMENTS

RESULT 1
AX589722/c
LOCUS AX589722 36 bp DNA linear PAT 24-JAN-2003
DEFINITION Sequence 29 from Patent WO02081621.
ACCESSION AX589722
VERSION AX589722.1 GI:27901012
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE 1
AUTHORS Loosmore,S.M. and Audonnet,J.C.
TITLE Vaccine against the nile fever virus
JOURNAL Patent: WO 02081621-A 29 17-OCT-2002;
MATERIAL (FR)
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					0;
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LOCUS	AX589721				
DEFINITION	Sequence 28 from Patent WO02081621.				
ACCESSION	AX589721				
VERSION	AX589721.1 GI:27901011				
KEYWORDS	synthetic construct				
SOURCE	other sequences; artificial sequences.				
ORGANISM					

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REFERENCE
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AUTHORS      Loomore,S.M. and Audonnet,J.C.
TITLE        Vaccine against the nile fever virus
JOURNAL      MENTAL (FR)
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1 TCATGACTGCAATTCGGTC 20
|||||
34 TCATGACTGCAATTCGGTC 15
Db
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LOCUS          AX589711 33 bp DNA linear PAT 24-JAN-2003
DEFINITION     Sequence 18 from Patent WO02081621.
ACCESSION      AX589711
VERSION        AX589711.1 GI:27901001
KEYWORDS       .
SOURCE         synthetic construct
ORGANISM       other sequences; artificial sequences.
REFERENCE
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AUTHORS      Loomore,S.M. and Audonnet,J.C.
TITLE        Vaccine against the nile fever virus
JOURNAL      MENTAL (FR)
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/db_xref="taxon:32630"
/note="oligonucleotide"
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Best Local Similarity 100.0%; Pred. No. 0.81;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY
1 TCATGACTGCAATTCGGT 19
|||||
28 TCATGACTGCAATTCGGT 10
Db
RESULT 4
AX589702/c
LOCUS          AX589702 36 bp DNA linear PAT 24-JAN-2003
DEFINITION     Sequence 9 from Patent WO02081621.
ACCESSION      AX589702
VERSION        AX589702.1 GI:27900992
KEYWORDS       .
SOURCE         synthetic construct
ORGANISM       other sequences; artificial sequences.
REFERENCE
1
AUTHORS      Loomore,S.M. and Audonnet,J.C.
TITLE        Vaccine against the nile fever virus
JOURNAL      MENTAL (FR)
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/db_xref="taxon:32630"
/note="oligonucleotide"
ORIGIN
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Best Local Similarity 100.0%; Pred. No. 0.81;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY
1 TCATGACTGCAATTCGGT 19
|||||
28 TCATGACTGCAATTCGGT 10
Db
RESULT 5
AX224249/c
LOCUS          AX224249 56 bp DNA linear PAT 10-SEP-2001
DEFINITION     Sequence 41 from Patent WO0160847.
ACCESSION      AX224249
VERSION        AX224249.1 GI:15554499
KEYWORDS       .
SOURCE         synthetic construct
ORGANISM       other sequences; artificial sequences.
REFERENCE
1
AUTHORS      Kinney,R.M., Kinney,C.Y., Butrapet,S., Gubler,D.L. and
Bhamarapravati,N.
TITLE        Avirulent, immunogenic flavivirus chimeras
JOURNAL      Patent: WO 0160847-A 41 23-AUG-2001;
The Secretary, Department of Health and Human Services (US)
FEATURES
source
1. .56
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1 TCATGACTGCAATTCGGT 19
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43 TCATGACTGCAATTCGGT 25
Db
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AX182204
LOCUS          AX182204 25 bp DNA linear PAT 06-AUG-2001
DEFINITION     Sequence 14 from Patent WO0142441.
ACCESSION      AX182204
VERSION        AX182204.1 GI:15133479
KEYWORDS       .
SOURCE         synthetic construct
ORGANISM       other sequences; artificial sequences.
REFERENCE
1
AUTHORS      Reddy,S.I., Sadhu,L.I., Shukla,V.C. and Ferraiolo,G.I.
TITLE        Plastid transformation
JOURNAL      Patent: WO 0142441-A 14 14-JUN-2001;
International Centre for Genetic Engineering and Biotechnology (IT)
FEATURES
source
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/mol_type="unassigned DNA"
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Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY
3 ATGACTGCAATT 14
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Qy 1 TCATGACTGCAA 12
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Db 9 TCATGACTGCAA 20

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RESULT 9					
AR444320					
LOCUS	AR444320		51 bp	DNA	
DEFINITION	Sequence 731 from patent US 6670464.			linear	PAT 20-FEB-2004
ACCESSION	AR444320				
VERSION	AR444320.1		GI:42672099		

REFERENCE	1. (bases 1 to 31)
AUTHORS	Shimkets, R. A. and Leach, M.
TITLE	Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
JOURNAL	Patent: US 6670464-A 731 30-DEC-2003;
FEATURES	Location/Qualifiers
source	1. 51

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Query Match          52.2%;   Score 12;   DB 6;   Length 51;
Best Local Similarity 100.0%;   Pred. No. 1.7e+04;
Matches 12;   Conservative 0;   Mismatches 0;   Indels 0;   Gaps 0;
Qy      5   GACTGCAATTC 16
Db      12  GACTGCAATTC 23

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SOURCE	UNKNOWN.
ORGANISM	Unknown.
REFERENCE	Unclassified.
AUTHORS	1 (bases 1 to 51)
TITLE	Shinketsu, R.A. and Leach, M. Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
JOURNAL	Patent: US 6670464-A 732 30-DEC-2003;
FEATURES	Location/Qualifiers 1..51 source

Qy 5 GACTGCAATTCC 16
|||
Db 12 GACTGCAATTCC 23

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RESULT 11
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DEFINITION Sequence 5678 from Patent W00210449.
ACCESSION CQ536043
VERSION CQ536043
KEYWORDS CQ536043.1 GI:41502307
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE Eukaryotic; Metazoa; Chordata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
TITLE Shoshan,A., Wasserman,A., Mintz,E., Mintz,L. and Faigler,S.
JOURNAL Oligonucleotide library for detecting rna transcripts and splice
FEATURES variants that populate a transcriptome
source Patent: WO 0210449-A 5678 07-FEB-2002;
Compugen Inc. (US)
LOCATION/Qualifiers
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Db 30 GACTGCAATTCC 41
RESULT 12
POLDIPI/c
LOCUS
DEFINITION Poliovirus defective interfering particle 17 mRNA, partial cds.
ACCESSION M30219
VERSION M30219.1 GI:332915
KEYWORDS Poliovirus
SOURCE Poliovirus
ORGANISM Viruses; ssRNA positive-strand viruses, no DNA stage;
REFERENCE Picornaviridae; Enterovirus.
AUTHORS Kuge,S., Saito,I. and Nomoto,A.
TITLE Primary structure of poliovirus defective-interfering particle
JOURNAL genomes and possible generation mechanisms of the particles
MEDLINE J. Mol. Biol. 192 (3), 473-487 (1986)
PUBMED 87169734
COMMENT Original source text: Poliovirus defective interfering particle 17,
CDNA to viral RNA.
FEATURES Location/Qualifiers
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Best Local Similarity 100.0%; Pred. No. 1.7e+04;
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QY 5 GACTGCAATTCC 16
Db 33 GACTGCAATTCC 22
RESULT 13
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LOCUS
DEFINITION Sequence 26 from patent US 6153382.
ACCESSION AR119503
VERSION AR119503.1 GI:14102202
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 15)
AUTHORS Karn,J., Gait,M.John., Heaphy,S. and Dingwall,C.
TITLE Viral growth inhibition
JOURNAL Patent: US 6153382-A 26 28-NOV-2000;
FEATURES Location/Qualifiers
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DEFINITION Sequence 8 from patent US 6528640.
ACCESSION AR285636
VERSION AR285636.1 GI:29723230
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 16)
AUTHORS Beigelman,L., Burgin,A., Beaudry,A., Karpeisky,A.,
TITLE Matulic-Adamic,J., Sweedler,D. and Zinnen,S.
JOURNAL Synthetic ribonucleic acids with RNase activity
FEATURES Patent: US 6528640-A 8 04-MAR-2003;
Location/Qualifiers
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Best Local Similarity 100.0%; Pred. No. 7.8e+04;
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AR397627/c
LOCUS
DEFINITION Sequence 8 from patent US 6617438.
ACCESSION AR397627
VERSION AR397627.1 GI:40134758
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 16)
AUTHORS Beigelman,L., Burgin,A.B., Beaudry,A., Karpeisky,A.,
TITLE Matulic-Adamic,J., Sweedler,D. and Zinnen,S.
JOURNAL Oligoribonucleotides with enzymatic activity
FEATURES Patent: US 6617438-A 8 09-SEP-2003;
Location/Qualifiers
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source
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DEFINITION Sequence 26 from patent US 6153382.
ACCESSION AR119503
VERSION AR119503.1 GI:14102202
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 15)
AUTHORS Karn,J., Gait,M.John., Heaphy,S. and Dingwall,C.
TITLE Viral growth inhibition
JOURNAL Patent: US 6153382-A 26 28-NOV-2000;
FEATURES Location/Qualifiers
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Best Local Similarity 100.0%; Pred. No. 7.8e+04;
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Db 14 AATTCCGGTCT 4
RESULT 14
AR285636/c
LOCUS
DEFINITION Sequence 8 from patent US 6528640.
ACCESSION AR285636
VERSION AR285636.1 GI:29723230
KEYWORDS
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ORGANISM
REFERENCE 1 (bases 1 to 16)
AUTHORS Beigelman,L., Burgin,A., Beaudry,A., Karpeisky,A.,
TITLE Matulic-Adamic,J., Sweedler,D. and Zinnen,S.
JOURNAL Synthetic ribonucleic acids with RNase activity
FEATURES Patent: US 6528640-A 8 04-MAR-2003;
Location/Qualifiers
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Db 13 GCAATTCGGT 3
RESULT 15
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LOCUS
DEFINITION Sequence 8 from patent US 6617438.
ACCESSION AR397627
VERSION AR397627.1 GI:40134758
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 16)
AUTHORS Beigelman,L., Burgin,A.B., Beaudry,A., Karpeisky,A.,
TITLE Matulic-Adamic,J., Sweedler,D. and Zinnen,S.
JOURNAL Oligoribonucleotides with enzymatic activity
FEATURES Patent: US 6617438-A 8 09-SEP-2003;
Location/Qualifiers
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ORIGIN

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